

```

DEFINITION HS_5142_A1_D07_SP6E_RPCI-11 Human Male BAC Library Homo sapiens
genomic clone Plate-718 Col-13 Row-G, DNA sequence.
ACCESSION AQ457751
VERSION AQ457751.1 GI:4636391
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 454)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallaceu.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 718 row: G column: 13
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 454.
FEATURES
source
Location/Qualifiers
1..454
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=718 Col=13 Row=G"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"
BASE COUNT 128 a 83 c 95 g 145 t 3 others
ORIGIN

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Query Match      85.6%; Score 15.4; DB 97; Length 454;
Best Local Similarity 94.1%; Pred. No. 8.4e+02;
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ctcccagcgtgcgcac 18
|||||
Db 300 CTCCCAGCGTGAGCCAT 316

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Search completed: December 4, 2000, 21:06:39
Job time: 19239 sec

Email: Robert.Strausberg@nih.gov
 This clone is available royalty-free through LLNL ; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Seq primer: -40UP from Gibco
 High quality sequence stop: 456.

FEATURES

Location/Qualifiers
 1. .525
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:2662007"
 /clone_lib="Soares_NFL_T_GBC_S1"
 /lab_host="DH10B"
 /note="Organ: pooled; Vector: pT73D-Pac (Pharmacia) with
 a modified polylinker; Site_1: Not 1; Site_2: Eco RI;
 Equal amounts of plasmid DNA from three normalized
 libraries (fetal lung NbHL19W, testis NHT, and B-cell
 NCI-CGAP.GC81) were mixed, and ss circles were made in
 vitro. Following HAP purification, this DNA was used as
 tracer in a subtractive hybridization reaction. The driver
 was PCR-amplified cDNAs from pools of 5,000 clones made
 from the same 3 libraries. The pools consisted of
 I.M.A.G.E. clones 297480-302087, 682632-687239,
 726408-728711, and 729096-731399. Subtraction by Bento
 Soares and M. Fatima Bonaldo."
 154 a 104 c 111 g 155 t 1 others

BASE COUNT

ORIGIN

Query Match 88.9%; Score 16; DB 20; Length 525;
 Best Local Similarity 100.0%; Pred. No. 4.3e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tccacgctgcgccat 18

Db 305 TCCACGCTGCGCCAT 320

RESULT 13

AI982970
 LOCUS wt46b05.x1 NCI-CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2510481 3'
 DEFINITION similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD PROTEIN. ;, mRNA
 sequence.
 AI982970
 VERSION AI982970.1 GI:5810189
 KEYWORDS EST.
 SOURCE human.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 735)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)

JOURNAL

COMMENT
 Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert.Strausberg@nih.gov
 Life Technologies catalog #: 11548-013
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 www-bio.llnl.gov/bbrp/image/image.html
 Seq primer: -40UP from Gibco
 High quality sequence stop: 426.

FEATURES

Location/Qualifiers
 1. .735
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:2510481"
 /clone_lib="NCI-CGAP_Pan1"
 /tissue_type="adenocarcinoma"
 /lab_host="DH10B"

/note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
 Site_2: NotI; Cloned unidirectionally. Primer: Oligo dT.
 Average insert size 1.72 kb. Life Technologies catalog #:
 11548-013"

BASE COUNT 215 a 150 c 153 g 217 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 735;
 Best Local Similarity 100.0%; Pred. No. 4.3e+02;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tccacgctgcgccat 18

Db 304 TCCACGCTGCGCCAT 319

RESULT 14

AQ214736
 LOCUS HS_3250_B1_F05_MR CIT Approved Human Genomic Sperm Library D Homo
 DEFINITION sapiens genomic clone Plate=3250 Col=9 Row=L, DNA sequence.
 AQ214736
 ACCESSION AQ214736.1 GI:3625937
 VERSION GSS.
 KEYWORDS SOURCE human.

ORGANISM

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 416)
 Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
 Hood,L.
 Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome
 Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
 99380589

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Sequence Tagged Connector
 Plate: 3250 row: L column: 9
 Class: BAC ends
 High quality sequence stop: 416.

FEATURES

source

Location/Qualifiers
 1. .416
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Plate=3250 Col=9 Row=L"
 /clone_lib="CIT Approved Human Genomic Sperm Library D"
 /sex="male"
 /note="Organ: sperm; Vector: pBelobAC11; BAC Clones in
 E-Coli DH10B"

BASE COUNT 120 a 123 c 72 g 100 t 1 others

ORIGIN

Query Match 85.6%; Score 15.4; DB 90; Length 416;
 Best Local Similarity 94.1%; Pred. No. 8.3e+02;
 Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 ctcccagctgcgccat 18

Db 177 CTACCAGCTGCGCCAT 193

RESULT 15

AQ457751

LOCUS

AQ457751 454 bp DNA GSS 23-APR-1999

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BASE COUNT      121 a      85 c      95 g      137 t      1 others
ORIGIN

Query Match      88.9%; Score 16; DB 19; Length 439;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tccacgctgcgccat 18
|||||
Db 306 TCCACGCTGC GCCAT 321

RESULT 10
AI005163
LOCUS      ou13c07.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
DEFINITION IMAGE:1626156 3' similar to TR:Q99499 Q99499 DYNEIN-RELATED
PROTEIN. ; mRNA sequence.
ACCESSION AI005163.1 GI:3214673
VERSION    1
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 439)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL   Unpublished (1997)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Tel: (301) 496-1550
            Email: Robert.Strausberg@nih.gov
            This clone is available royalty-free through LLNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            Insert Length: 703 Std Error: 0.00
            Seq primer: primer name ambiguous
            High quality sequence stop: 444.
            Location/Qualifiers
                location=1..459
                organism="Homo sapiens"
                db_xref="taxon:9606"
                clone="IMAGE:1626156"
                clone_lib="Soares_NFL_T_GBC_S1"
                lab_host="DH10B"
                note="Organ: pooled; Vector: pT7T3D-Pac (Pharmacia) with
                a modified polylinker; Site_1: Not I; Site_2: Eco RI;
                Equal amounts of plasmid DNA from three normalized
                libraries (fetal lung NBHL19W, testis NHT, and B-cell
                NCI-CGAP-GCB1) were mixed, and ss circles were made in
                vitro. Following HAP purification, this DNA was used as
                tracer in a subtractive hybridization reaction. The driver
                was PCR-amplified cDNAs from pools of 5,000 clones made
                from the same 3 libraries. The pools consisted of
                I.M.A.G.E. clones 297480-302087, 682632-687239,
                726408-728711, and 729096-731399. Subtraction by Bento
                Soares and M. Fatima Bonaldo.
BASE COUNT      128 a      83 c      96 g      152 t
ORIGIN

Query Match      88.9%; Score 16; DB 7; Length 459;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tccacgctgcgccat 18
|||||
Db 350 TCCACGCTGC GCCAT 365

RESULT 11

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AI682599      510 bp      mRNA      EST      17-DEC-1999
LOCUS      wc63e11.x1 NCI_CGAP_Panl Homo sapiens cDNA clone IMAGE:2323340 3'
DEFINITION similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD PROTEIN. ; mRNA
            sequence.
ACCESSION AI682599.1 GI:4892781
VERSION    1
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 510)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL   Unpublished (1997)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Tel: (301) 496-1550
            Email: Robert.Strausberg@nih.gov
            Life Technologies catalog #: 11548-013
            DNA sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www-bio.llnl.gov/bbrp/image/image.html
            Insert Length: 1483 Std Error: 0.00
            Seq primer: -40UP from Gibco
            High quality sequence stop: 360.
            Location/Qualifiers
                location=1..510
                organism="Homo sapiens"
                db_xref="taxon:9606"
                clone="IMAGE:2323340"
                clone_lib="NCI_CGAP_Panl"
                tissue_type="adenocarcinoma"
                lab_host="DH10B"
                note="Organ: pancreas; Vector: pCMV-SPORT6; Site_1: SalI;
                Site_2: NotI; Cloned unidirectionally. Primer: Oligo dt.
                Average insert size 1.72 kb. Life Technologies catalog #:
                11548-013"
BASE COUNT      143 a      89 c      110 g      168 t
ORIGIN

Query Match      88.9%; Score 16; DB 12; Length 510;
Best Local Similarity 100.0%; Pred. No. 4.2e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 tccacgctgcgccat 18
|||||
Db 349 TCCACGCTGC GCCAT 364

RESULT 12
AI182837
LOCUS      xj64d12.x1 Soares_NFL_T_GBC_S1 Homo sapiens cDNA clone
DEFINITION IMAGE:2662007 3' similar to TR:O43326 O43326 HYPOTHETICAL 65.4 KD
            PROTEIN. ; mRNA sequence.
ACCESSION AI182837
VERSION    1
KEYWORDS   EST.
SOURCE     human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 525)
AUTHORS   NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE     National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
JOURNAL   Unpublished (1997)
COMMENT   Contact: Robert Strausberg, Ph.D.
            Tel: (301) 496-1550

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RESULT 7
AII08632      658 bp      mRNA      EST      18-NOV-1998
LOCUS      GH07838.5prime GH Drosophila melanogaster head pOT2 Drosophila
DEFINITION      melanogaster cDNA clone GH07838 5prime, mRNA sequence.
ACCESSION      AII08632
VERSION      AII08632
KEYWORDS      EST.
SOURCE      AII08632.1 GI:3477167
            fruit fly.
ORGANISM      Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE      1 (bases 1 to 658)
AUTHORS      Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein
            ,P., Lewis,S. and Rubin,G.M.
TITLE      BDGP/HMI Drosophila EST Project
JOURNAL      Unpublished (1997)
COMMENT      Contact: Harvey, D.
            G. M. Rubin-Molecular and Cell Biology
            University of California Berkeley
            539 LSA, Berkeley, CA 94720-3200, USA
            Fax: 510 643 9947
            Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
            Plate: 78 row: D column: 2
            High quality sequence stop: 520.
FEATURES      Location/Qualifiers
            1..658
            /organism="Drosophila melanogaster"
            /db_xref="taxon:7227"
            /clone_lib="GH07838"
            /sex="male and female"
            /dev_stage="adult"
            /lab_host="DH5 - alpha"
            /note="Organ: head; Vector: pOT2; Site_1: EcoRI; Site_2:
            XhoI; Sized fractionated cDNAs were directly ligated into
            pOT2. Plasmid cDNA library."
BASE COUNT      167 a 188 c 190 g 113 t
ORIGIN
1..658
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_lib="GH07838"
/sex="male and female"
/dev_stage="adult"
/lab_host="DH5 - alpha"
/note="Organ: head; Vector: pOT2; Site_1: EcoRI; Site_2:
XhoI; Sized fractionated cDNAs were directly ligated into
pOT2. Plasmid cDNA library."
Query Match      91.1%; Score 16.4; DB 8; Length 658;
Best Local Similarity 94.4%; Pred. NO. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
|| |||||
Db 80 TCGCCACGCGTGCACAT 97

RESULT 8
AA990787      695 bp      mRNA      EST      24-NOV-1998
LOCUS      LD34664.5prime LD Drosophila melanogaster embryo pOT2 Drosophila
DEFINITION      melanogaster cDNA clone LD34664 5prime, mRNA sequence.
ACCESSION      AA990787
VERSION      AA990787
KEYWORDS      EST.
SOURCE      AA990787.1 GI:3177320
            fruit fly.
ORGANISM      Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE      1 (bases 1 to 695)
AUTHORS      Harvey,D., Hong,L., Evans-Holm,M., Pendleton,J., Su,C., Brokstein
            ,P., Lewis,S. and Rubin,G.M.
TITLE      BDGP/HMI Drosophila EST Project
JOURNAL      Unpublished (1997)
COMMENT      Contact: Harvey, D.
            G. M. Rubin-Molecular and Cell Biology
            University of California Berkeley
            539 LSA, Berkeley, CA 94720-3200, USA
            Fax: 510 643 9947
            Email: http://www.fruitfly.org/EST, est@fruitfly.berkeley.edu
            Plate: 78 row: D column: 2
            High quality sequence stop: 520.
FEATURES      Location/Qualifiers
            1..695
            /organism="Drosophila melanogaster"
            /db_xref="taxon:7227"
            /clone_lib="LD34664"
            /sex="male and female"
            /dev_stage="embryo"
            /lab_host="XLI Blue"
            /note="Organ: embryo; Vector: pOT2; Site_1: EcoRI; Site_2:
            XhoI; Sized fractionated cDNAs were directly ligated into
            pOT2."
BASE COUNT      178 a 201 c 206 g 110 t
ORIGIN
1..695
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone_lib="LD34664"
/sex="male and female"
/dev_stage="embryo"
/lab_host="XLI Blue"
/note="Organ: embryo; Vector: pOT2; Site_1: EcoRI; Site_2:
XhoI; Sized fractionated cDNAs were directly ligated into
pOT2."
Query Match      91.1%; Score 16.4; DB 7; Length 695;
Best Local Similarity 94.4%; Pred. NO. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
|| |||||
Db 66 TCGCCACGCGTGCACAT 83

RESULT 9
AW070294      439 bp      mRNA      EST      13-OCT-1999
LOCUS      xa06g11.x1 Soares_NFL_T.GBC.S1 Homo sapiens cDNA clone
DEFINITION      IMAGE:2567588 3' similar to TR:043326 O43326 HYPOTHETICAL 65.4 KD
            PROTEIN: ;, mRNA sequence.
ACCESSION      AW070294
VERSION      AW070294.1 GI:6025292
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 439)
AUTHORS      NCI-CCAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
JOURNAL      Contact: Robert Strausberg, Ph.D.
COMMENT      Tel: (301) 496-1550
            Email: Robert_Strausberg@nih.gov
            This clone is available royalty-free through LML; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            Seq primer: -400P from Gibco
            High quality sequence stop: 400.
FEATURES      Location/Qualifiers
            1..439
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /clone_lib="IMAG2567588"
            /lab_host="Soares_NFL_T.GBC.S1"
            /note="Organ: pooled; Vector: pT73p-Pac (Pharmacia) with
            a modified polylinker; Site_1: Not I; Site_2: Eco RI;
            Equal amounts of plasmid DNA from three normalized
            libraries (fetal lung NBHL19W, testis NHT, and B-cell
            NCI-CGAP-GC81) were mixed, and ss circles were made in
            vitro. Following HAP purification, this DNA was used as
            tracer in a subtractive hybridization reaction. The driver
            was PCR-amplified cDNAs from pools of 5,000 clones made
            from the same 3 libraries. The pools consisted of
            I.M.A.G.E. clones 297480-302087, 682632-687239,
            726408-728711, and 729096-731399. Subtraction by Bento

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VERSION      H83468.1  GI:1062139
KEYWORDS     EST.
SOURCE       human.
ORGANISM     Homo sapiens
REFERENCE    1 (bases 1 to 398)
AUTHORS      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman
              , M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
              Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston
              , R., Williamson, A., Wohldmann, P. and Wilson, R.
              The WashU-Merck EST Project
              Unpublished (1995)
              Contact: Wilson RK
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
              Tel: 314 286 1800
              Fax: 314 286 1810
              Email: est@watson.wustl.edu
              High quality sequence stop: 305
              Source: IMAGE Consortium, LLNL
              This clone is available royalty-free through LLNL; contact the
              IMAGE Consortium (info@image.llnl.gov) for further information.
              Insert Length: 1825 Std Error: 0.00
              Seq primer: M13Rp1
              High quality sequence stop: 305.
FEATURES     Location/Qualifiers
              1..398
               /organism="Homo sapiens"
               /db_xref="GDB:3850898"
               /db_xref="taxon:9606"
               /clone="IMAGE:222137"
               /clone_lib="Soares retina N2b5HR"
               /sex="male"
               /tissue_type="retina"
               /dev_stage="55 year old"
               /lab_host="DH10B (ampicillin resistant)"
               /note="Organ: eye; Vector: pT7T3D (Pharmacia) with a
               modified polylinker; Site.1: Not I; Site.2: Eco RI; 1st
               strand cDNA was primed with a Not I - oligo(dT) primer [5'
               TGTACCAATCTGAAGTCGAGCGCGCGCTTTTCTTTTCTTTT 3'],
               double-stranded cDNA was size selected, ligated to Eco RI
               adapters (Pharmacia), digested with Not I and cloned into
               the Not I and Eco RI sites of a modified pT7T3 vector
               (Pharmacia). The retinas were obtained from a 55 year old
               Caucasian and total cellular poly(A)+ RNA was extracted 6
               hrs after their removal. The retina RNA was kindly
               provided by Roderick R. McInnes M.D. Ph.D. from the
               University of Toronto. Library constructed by Bento
               Soares and M.Fatima Bonaldo."
BASE COUNT   85 a 104 c 108 g 98 t 3 others
ORIGIN

Query Match 91.1%; Score 16.4; DB 37; Length 398;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
    |||||
Db 324 TCTCCACGCGTGGGCCAT 307

RESULT      3
LOCUS       AA402089
DEFINITION  AA402089 412 bp mRNA EST 16-MAY-1997
IMAGE:741723 5' similar to SW:YK18_YEAST P36132 HYPOTHETICAL 46.6
KD PROTEIN IN DAL80-GAP1 INTERGENIC REGION. ;, mRNA sequence.
ACCESSION   AA402089
VERSION     AA402089.1 GI:2056072
KEYWORDS     EST.
SOURCE      human.

ORGANISM     Homo sapiens
REFERENCE    1 (bases 1 to 442)
AUTHORS      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman
              , M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
              Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston
              , R., Williamson, A., Wohldmann, P. and Wilson, R.
              The WashU-Merck EST Project
              Unpublished (1995)
              Contact: Wilson RK
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
              Tel: 314 286 1800
              Fax: 314 286 1810
              Email: est@watson.wustl.edu
              High quality sequence stop: 378.
              Source: IMAGE Consortium, LLNL
              This clone is available royalty-free through LLNL; contact the
              IMAGE Consortium (info@image.llnl.gov) for further information.
              Seq primer: -28m13 rev2 ET from Amersham
              High quality sequence stop: 378.
FEATURES     Location/Qualifiers
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               /organism="Homo sapiens"
               /db_xref="GDB:5941912"
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               /clone="IMAGE:741723"
               /clone_lib="Soares ovary tumor N8HOT"
               /sex="female"
               /tissue_type="ovarian tumor"
               /lab_host="DH10B (ampicillin resistant)"
               /note="Organ: ovary; Vector: pT7T3D (Pharmacia) with a
               modified polylinker; Site.1: Not I; Site.2: Eco RI; 1st
               strand cDNA was primed with a Not I - oligo(dT) primer [5'
               TGTACCAATCTGAAGTCGAGCGCGCGCTTTTCTTTTCTTTT 3'],
               double-stranded cDNA was size selected, ligated to Eco RI
               adapters (Pharmacia), digested with Not I and cloned into
               the Not I and Eco RI sites of a modified pT7T3 vector
               (Pharmacia). Library constructed by Bento Soares and
               M.Fatima Bonaldo."
BASE COUNT   77 a 115 c 131 g 89 t
ORIGIN

Query Match 91.1%; Score 16.4; DB 4; Length 412;
Best Local Similarity 94.4%; Pred. No. 2.7e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
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Db 77 TCTCCACGCGTGGGCCAT 94

RESULT      4
LOCUS       W96258/c
DEFINITION  ze42b06.r1 Soares retina N2b4HR Homo sapiens cDNA clone
IMAGE:361619 5', mRNA sequence.
ACCESSION   W96258
VERSION     W96258.1 GI:1426165
KEYWORDS     EST.
SOURCE      human.
ORGANISM     Homo sapiens
REFERENCE    1 (bases 1 to 442)
AUTHORS      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman
              , M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
              Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston
              , R., Williamson, A., Wohldmann, P. and Wilson, R.
              The WashU-Merck EST Project
              Unpublished (1995)
              Contact: Wilson RK
              Washington University School of Medicine
              4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
              Tel: 314 286 1800
              Fax: 314 286 1810
              Email: est@watson.wustl.edu
              High quality sequence stop: 378.
              Source: IMAGE Consortium, LLNL
              This clone is available royalty-free through LLNL; contact the
              IMAGE Consortium (info@image.llnl.gov) for further information.
              Seq primer: -28m13 rev2 ET from Amersham
              High quality sequence stop: 378.
FEATURES     Location/Qualifiers
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               /db_xref="GDB:5941912"
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               /clone="IMAGE:741723"
               /clone_lib="Soares ovary tumor N8HOT"
               /sex="female"
               /tissue_type="ovarian tumor"
               /lab_host="DH10B (ampicillin resistant)"
               /note="Organ: ovary; Vector: pT7T3D (Pharmacia) with a
               modified polylinker; Site.1: Not I; Site.2: Eco RI; 1st
               strand cDNA was primed with a Not I - oligo(dT) primer [5'
               TGTACCAATCTGAAGTCGAGCGCGCGCTTTTCTTTTCTTTT 3'],
               double-stranded cDNA was size selected, ligated to Eco RI
               adapters (Pharmacia), digested with Not I and cloned into
               the Not I and Eco RI sites of a modified pT7T3 vector
               (Pharmacia). Library constructed by Bento Soares and
               M.Fatima Bonaldo."
BASE COUNT   77 a 115 c 131 g 89 t
ORIGIN

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 15:46:00 ; Search time 1141.84 Seconds
(without alignments)
97.466 Million cell updates/sec

Title: US-09-369-941-1
Perfect score: 18
Sequence: 1 tctccagcgtgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 7189864 seqs, 3091403243 residues

Total number of hits satisfying chosen parameters: 14379728

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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2: gb_est2:*
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107: gb_gss16:*
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112: gb_gss21:*
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114: gb_gss23:*
115: gb_gss24:*
116: em_gss5:*

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM nucleic - nucleic search, using sw model
Run on: December 4, 2000, 15:46:00 ; Search time 993.06 Seconds
(without alignments)
79.166 Million cell updates/sec

Title: US-09-369-941-1
Perfect score: 18
Sequence: 1 tctcccagctgcgccat 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1033670 seqs, 2183789903 residues 2067340
Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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2: gb_ba2 :
3: gb_om :
4: gb_ov :
5: gb_pat :
6: gb_ph :
7: gb_pll :
8: gb_pl2 :
9: gb_pr1 :
10: gb_pr2 :
11: gb_pr3 :
12: gb_ro :
13: gb_sy :
14: gb_un :
15: em_fun :
16: em_hum1 :
17: em_hum2 :
18: em_in :
19: em_om :
20: em_or :
21: em_ov :
22: em_pat :
23: em_ph :
24: em_pl :
25: em_ro :
26: em_sts :
27: em_sy :
28: em_un :
29: em_vi :
30: gb_ba3 :
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32: gb_in2 :
33: gb_in3 :
34: gb_pl3 :
35: gb_pr4 :
36: em_ba1 :
37: em_ba2 :
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42: em_htg5 :
43: em_htg6 :

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45: em_htg8 :
46: em_htg9 :
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64: em_hum6 :
65: gb_pr5 :
66: gb_pr6 :
67: gb_pr7 :
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69: gb_htg2 :
70: gb_htg3 :
71: gb_htg4 :
72: gb_htg5 :
73: gb_htg6 :
74: gb_htg7 :
75: gb_htg8 :
76: gb_htg9 :
77: gb_htg10 :
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84: gb_htg17 :
85: gb_htg18 :
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87: gb_htg20 :
88: gb_htg21 :
89: gb_htg22 :
90: gb_htg23 :
91: gb_sts1 :
92: gb_sts2 :
93: gb_vil1 :
94: gb_vil2 :

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	18	100.0	18	5	AR052619	Sequence
2	18	100.0	18	5	AR052624	Sequence
3	18	100.0	18	5	196098	Sequence 17
c 4	18	100.0	27	5	AR004426	Sequence
c 5	18	100.0	27	5	143661	Sequence 13
c 6	18	100.0	27	5	186720	Sequence 8
c 7	18	100.0	35	5	AR052604	Sequence
c 8	18	100.0	35	5	196083	Sequence 2
c 9	18	100.0	454	67	S72602	bc12 [human
c 10	18	100.0	615	5	AR052623	Sequence
c 11	18	100.0	717	5	AR052622	Sequence
c 12	18	100.0	760	5	AR021160	Sequence

C 13 18 100.0 755 5 A76121
C 14 18 100.0 911 67 HUMBL2B
C 15 18 100.0 1846 5 AR054009
C 16 18 100.0 1846 66 HSBCL21G
C 17 18 100.0 5086 5 AR052621
C 18 18 100.0 5086 5 AR054008
C 19 18 100.0 5086 5 HUMBL2A
C 20 18 100.0 5105 5 I08038
C 21 18 100.0 6030 67 HUMBL2C
C 22 18 100.0 86692 75 AC021803
C 23 18 100.0 169542 89 AP001915
C 24 18 100.0 188601 69 AC009267
C 25 17 94.4 17 5 I96090
C 26 17 94.4 35737 74 AC005263
C 27 16.4 91.1 67568 79 AC020378
C 28 16.4 91.1 99614 68 AC008358
C 29 16.4 91.1 128132 68 AC007732
C 30 16.4 91.1 161799 9 AC002091
C 31 16.4 91.1 170757 68 AC007608
C 32 16.4 91.1 186323 68 AC006491
C 33 16.4 91.1 210299 31 AE003690
C 34 16.4 91.1 217026 72 AC013669
C 35 16 88.9 157432 83 AC069438
C 36 16 88.9 158934 81 AC055780
C 37 16 88.9 211341 85 AL138898
C 38 15.4 85.6 10854 1 AE001886
C 39 15.4 85.6 38616 9 AC004208
C 40 15.4 85.6 42245 9 AC004476
C 41 15.4 85.6 75163 83 AC069103
C 42 15.4 85.6 75533 68 AC006412
C 43 15.4 85.6 112630 78 AC025461
C 44 15.4 85.6 117143 67 HUMBL2C
C 45 15.4 85.6 120841 10 AC008162

ALIGNMENTS

RESULT 1
LOCUS AR052619 18 bp DNA
DEFINITION Sequence 17 from patent US 5831066.
ACCESSION AR052619
VERSION AR052619.1 GI:5975983
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Reed,J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 17 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..18
BASE COUNT 2 a 8 c 4 g 4 t
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Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 tctccagcgtgcgcacat 18
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Db 1 TCTCCAGCGTGCACAT 18
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LOCUS AR052624 18 bp DNA
DEFINITION Sequence 24 from patent US 5831066.
ACCESSION AR052624
VERSION AR052624.1 GI:5975983
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Reed,J.C. and Sato,T.
TITLE Nucleic acids encoding Fas associated proteins and screening assays
JOURNAL Patent: US 5747245-A 13 05-MAY-1998;
FEATURES Location/Qualifiers
source 1..27
BASE COUNT 2 a 8 c 4 g 4 t
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Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 TCTCCAGCGTGCACAT 18
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LOCUS AR052619 18 bp DNA
DEFINITION Sequence 17 from patent US 5734033.
ACCESSION AR052619
VERSION AR052619.1 GI:3940568
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 18)
AUTHORS Reed,J.C.
TITLE Antisense oligonucleotides inhibiting human bcl-2 gene expression -
JOURNAL Patent: US 5734033-A 17 31-MAR-1998;
FEATURES Location/Qualifiers
source 1..18
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Db 1 TCTCCAGCGTGCACAT 18
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LOCUS AR004426/c 27 bp DNA
DEFINITION Sequence 13 from patent US 5747245.
ACCESSION AR004426
VERSION AR004426.1 GI:3965305
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Reed,J.C. and Sato,T.
TITLE Nucleic acids encoding Fas associated proteins and screening assays
JOURNAL Patent: US 5747245-A 13 05-MAY-1998;
FEATURES Location/Qualifiers
source 1..27
BASE COUNT 2 a 8 c 4 g 4 t
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Best Local Similarity 100.0%; Pred. No. 83;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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|||||
Db 1 TCTCCAGCGTGCACAT 18
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DEFINITION Sequence 13 from patent US 5747245.
ACCESSION AR004426
VERSION AR004426.1 GI:3965305
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 27)
AUTHORS Reed,J.C. and Sato,T.
TITLE Nucleic acids encoding Fas associated proteins and screening assays
JOURNAL Patent: US 5747245-A 13 05-MAY-1998;
FEATURES Location/Qualifiers
source 1..27
BASE COUNT 2 a 8 c 4 g 4 t
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Best Local Similarity 100.0%; Pred. No. 83;
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|||||
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ORIGIN

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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 25 TCTCCCAGCGTGGCCCAT 8

RESULT 5
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LOCUS      I43661      27 bp      DNA      PAT      07-OCT-1997
DEFINITION Sequence 13 from patent US 5632994.
ACCESSION  I43661
VERSION     I43661.1 GI:2468759
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 27)
AUTHORS    Reed,J.C. and Sato,T.
TITLE      Fas associated proteins
JOURNAL    Patent: US 5632994-A 13 27-MAY-1997;
FEATURES   Location/Qualifiers
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BASE COUNT      7 a      6 c      10 g      4 t
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Qy 1 tctcccagcgtgcgccat 18
Db 25 TCTCCCAGCGTGGCCCAT 8

RESULT 6
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LOCUS      I86720      27 bp      DNA      PAT      10-JUN-1998
DEFINITION Sequence 8 from patent US 5702897.
ACCESSION  I86720
VERSION     I86720.1 GI:3206438
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 27)
AUTHORS    Reed,J.C. and Sato,T.
TITLE      Interaction of proteins involved in a cell death pathway
JOURNAL    Patent: US 5702897-A 8 30-DEC-1997;
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Qy 1 tctcccagcgtgcgccat 18
Db 25 TCTCCCAGCGTGGCCCAT 8

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RESULT 7
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LOCUS      AR052604      35 bp      DNA      PAT      29-SEP-1999
DEFINITION Sequence 2 from patent US 5831066.
ACCESSION  AR052604
VERSION     AR052604.1 GI:5975968
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 35)
AUTHORS    Reed,J.C.
TITLE      Regulation of bcl-2 gene expression
JOURNAL    Patent: US 5831066-A 2 03-NOV-1998;
FEATURES   Location/Qualifiers
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BASE COUNT      6 a      8 c      13 g      8 t
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Db 35 TCTCCCAGCGTGGCCCAT 18

RESULT 8
I96083/c
LOCUS      I96083      35 bp      DNA      PAT      01-DEC-1998
DEFINITION Sequence 2 from patent US 5734033.
ACCESSION  I96083
VERSION     I96083.1 GI:3940553
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 35)
AUTHORS    Reed,J.
TITLE      Antisense oligonucleotides inhibiting human bcl-2 gene expression
JOURNAL    Patent: US 5734033-A 2 31-MAR-1998;
FEATURES   Location/Qualifiers
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BASE COUNT      6 a      8 c      13 g      8 t
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Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
Db 35 TCTCCCAGCGTGGCCCAT 18

RESULT 9
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LOCUS      S72602      454 bp      DNA      PRI      07-MAY-1993
DEFINITION bcl2 [human, 697 pre-B cell acute lymphocytic leukemia cell line,
Genomic, 454 nt].
ACCESSION  S72602
VERSION     S72602.1 GI:241046
KEYWORDS   .
SOURCE     human 697 pre-B cell acute lymphocytic leukemia cell line.
ORGANISM   Homo sapiens
REFERENCE  1 (bases 1 to 454)

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AUTHORS Tanaka, S., Louie, D.C., Kant, J.A. and Reed, J.C.
TITLE Frequent incidence of somatic mutations in translocated BCL2
oncogenes of non-Hodgkin's lymphomas
JOURNAL Blood 79 (1), 229-237 (1992)
MEDLINE 92096610
REMARK GenBank staff at the National Library of Medicine created this
entry [NCBI gibbsq 72602] from the original journal article.
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/db_xref="taxon:9606"
gene 1..454
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CDS 41..433
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/translation="MAHAGRTGYDNRREIVMKYHYKLSQRYEWDAAGDVGAAAPGAP
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BASE COUNT 65 a 170 c 150 g 69 t
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Best Local Similarity 100.0%; Pred. No. 46;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
Db 58 TCTCCAGCGTGCACAT 41

RESULT 10
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LOCUS AR052623 615 bp DNA
DEFINITION Sequence 22 from patent US 5831066.
ACCESSION AR052623
VERSION AR052623.1 GI:5975987
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 615)
AUTHORS Reed, J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 22 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..615
/organism="unknown"
BASE COUNT 98 a 203 c 213 g 101 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 43;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
Db 18 TCTCCAGCGTGCACAT 1

RESULT 11
AR052622/c
LOCUS AR052622 717 bp DNA
DEFINITION Sequence 20 from patent US 5831066.
ACCESSION AR052622
VERSION AR052622.1 GI:5975986
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 717)
AUTHORS Reed, J.C.
TITLE Regulation of bcl-2 gene expression
JOURNAL Patent: US 5831066-A 20 03-NOV-1998;
FEATURES Location/Qualifiers
source 1..717
/organism="unknown"
BASE COUNT 113 a 237 c 237 g 130 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 717;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
Db 18 TCTCCAGCGTGCACAT 1

RESULT 12
AR021160/c
LOCUS AR021160 760 bp DNA
DEFINITION Sequence 11 from patent US 5789389.
ACCESSION AR021160
VERSION AR021160.1 GI:3975775
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 760)
AUTHORS Tarasiewicz, D.G., Schott, B., Holzmayer, T.A. and Robinson, I.B.
TITLE BCL2 derived genetic elements associated with sensitivity to
chemotherapeutic drugs
JOURNAL Patent: US 5789389-A 11 04-AUG-1998;
FEATURES Location/Qualifiers
source 1..760
/organism="unknown"
BASE COUNT 122 a 250 c 246 g 142 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 760;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgcac 18
Db 38 TCTCCAGCGTGCACAT 21

RESULT 13
A76121/c
LOCUS A76121 765 bp DNA
DEFINITION Sequence 1 from Patent WO9320200.
ACCESSION A76121
VERSION A76121.1 GI:6088257
KEYWORDS
SOURCE human.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 765)
AUTHORS Evan, G.I.
TITLE MODIFIED CELLS AND METHOD OF TREATMENT
JOURNAL Patent: WO 9320200-A 14-OCT-1993;
FEATURES IMP CANCER RES TECH (GB); EVAN GERARD IAN (GB)
Location/Qualifiers
source 1..765

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
31. .750
/note="unnamed protein product"
/codon_start=1
/protein_id="CAB8588.1"
/db_xref="GI:6088258"
/translation="MAHAGRTGYDNEIYMKYIHYKLSORGYENDAGDVGAAPPCAAP
APGIFSSQGHPIHPAASRDVARTSPLOTPAAGAAAGPALSPVPVPHALRQAGD
DFSRRYRGDAEMSSQLHTPTFARGFATVVEELFRDGVNMGWDAFEVFGVNCVE
SVNREMSPLVDNIALWMTEYLNRLHTWIQDNGWDAFEVFGVNCVE
TLLSLALVGCITLCAYLSHK" 144 t

BASE COUNT 120 a 251 c 250 g 144 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 765;
Best Local Similarity 100.0%; Pred. No. 42;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctccacagctgcgcacat 18
|||||
Db 48 TCTCCACGCTGCGCCAT 31

RESULT 14
HUMCL2B/c
LOCUS
DEFINITION
HUMAN B-cell leukemia/lymphoma 2 (bcl-2) proto-oncogene mRNA
encoding bcl-2-beta protein, complete cds.
ACCESSION
M13995
VERSION
M13995.1 GI:179368
KEYWORDS
alternative splicing; bcl-2-beta protein; proto-oncogene.
SOURCE
Human pre-B-cell leukemia cell line 380, cDNA to mRNA, clones
B[15,16]; and DNA, clone lambda-18-27.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 911)
Tsujiimoto, Y. and Croce, C.M.
Analysis of the structure, transcripts, and protein products of
bcl-2, the gene involved in human follicular lymphoma
Proc. Natl. Acad. Sci. U.S.A. 83 (14), 5214-5218 (1986)
86259760
COMMENT
Clean copy sequence for [1] kindly provided by Y. Tsujiimoto,
10-FEB-1987. The bcl-2 gene is transcribed by alternative splicing
into three mRNAs of different sizes. It consists of at least two
exons and encodes two proteins which only differ at their
carboxy-terminal ends, and it is activated by translocation into
proximity with the Ig heavy chain locus. Both the normal and
rearranged bcl-2 gene products are expressed in the B-cell
leukemia/lymphoma 2 cells. Genomic clone lambda-18-27 contained
all the DNA sequences on the 5' of the splice site (position 732).

FEATURES
Source
1. .911
/organism="Homo sapiens"
/db_xref="taxon:9606"
/map="18q21.3"
<1. .>911
/note="bcl2a mRNA"
147. .764
/gene="BCL2"
147. .764
/gene="BCL2"
/note="bcl2-beta protein"
/codon_start=1
/db_xref="GDB:G00-119-031"
/protein_id="AAA51814.1"
/db_xref="GI:179369"
/translation="MAHAGRTGYDNEIYMKYIHYKLSORGYENDAGDVGAAPPCAAP
APGIFSSQGHPIHPAASRDVARTSPLOTPAAGAAAGPALSPVPVPHALRQAGD
DFSRRYRGDAEMSSQLHTPTFARGFATVVEELFRDGVNMGWDAFEVFGVNCVE

misc_feature
/gene="BCL2"
732
/note="alternative splice donor (intron A start)"
BASE COUNT 156 a 281 c 306 g 168 t
ORIGIN 556 bp upstream of SstI site.

Query Match 100.0%; Score 18; DB 67; Length 911;
Best Local Similarity 100.0%; Pred. No. 40;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctccacagctgcgcacat 18
|||||
Db 164 TCTCCACGCTGCGCCAT 147

RESULT 15
AR054009/c
LOCUS
DEFINITION
Sequence 16 from patent US 5834306.
ACCESSION
AR054009
VERSION
AR054009.1 GI:5978871
KEYWORDS
Unknown.
SOURCE
Unknown.
ORGANISM
Unclassified.
REFERENCE
1 (bases 1 to 1846)
AUTHORS
Webster, K.A. and Bishopric, N.H.
TITLE
Tissue specific hypoxia regulated therapeutic constructs
JOURNAL
Patent: US 5834306-A 16 10-NOV-1998;
FEATURES
Location/Qualifiers
1. .1846
source
BASE COUNT 424 a 520 c 483 g 419 t
ORIGIN

Query Match 100.0%; Score 18; DB 5; Length 1846;
Best Local Similarity 100.0%; Pred. No. 36;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctccacagctgcgcacat 18
|||||
Db 904 TCTCCACGCTGCGCCAT 887

Search completed: December 4, 2000, 20:47:25
Job time: 18085 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 16:32:02 ; Search time 99.31 Seconds
(without alignments)
68.089 Million cell updates/sec

Title: US-09-369-941-1

Perfect score: 18
Sequence: 1 tctcccagcgtgcgcatt 18

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 480022 seqs, 187831343 residues

Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_36:*
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2: /cgn2_2/gcgdata/geneseq/geneseq/NA1981.DAT:*
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4: /cgn2_2/gcgdata/geneseq/geneseq/NA1983.DAT:*
5: /cgn2_2/gcgdata/geneseq/geneseq/NA1984.DAT:*
6: /cgn2_2/gcgdata/geneseq/geneseq/NA1985.DAT:*
7: /cgn2_2/gcgdata/geneseq/geneseq/NA1986.DAT:*
8: /cgn2_2/gcgdata/geneseq/geneseq/NA1987.DAT:*
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11: /cgn2_2/gcgdata/geneseq/geneseq/NA1990.DAT:*
12: /cgn2_2/gcgdata/geneseq/geneseq/NA1991.DAT:*
13: /cgn2_2/gcgdata/geneseq/geneseq/NA1992.DAT:*
14: /cgn2_2/gcgdata/geneseq/geneseq/NA1993.DAT:*
15: /cgn2_2/gcgdata/geneseq/geneseq/NA1994.DAT:*
16: /cgn2_2/gcgdata/geneseq/geneseq/NA1995.DAT:*
17: /cgn2_2/gcgdata/geneseq/geneseq/NA1996.DAT:*
18: /cgn2_2/gcgdata/geneseq/geneseq/NA1997.DAT:*
19: /cgn2_2/gcgdata/geneseq/geneseq/NA1998.DAT:*
20: /cgn2_2/gcgdata/geneseq/geneseq/NA1999.DAT:*
21: /cgn2_2/gcgdata/geneseq/geneseq/NA2000.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	16 Q86659	Bcl-2 antisense oligonucleotide
2	18	100.0	18	19 V52545	Unmethylated CpG d
3	18	100.0	18	19 V27719	Immunostimulatory
4	18	100.0	18	19 V28181	Antisense oligonucleotide
5	18	100.0	18	19 V19667	Human bcl-2 antisense
6	18	100.0	18	20 Z31944	CpG adjuvant oligo
7	18	100.0	18	20 Z41905	IL-12 secretion in
8	18	100.0	18	20 Z41948	IL-12 secretion in
9	18	100.0	18	20 X78803	HPV fusion protein
10	18	100.0	18	20 X88537	Cytosine-guanosine
11	18	100.0	18	20 X33514	BCL2-targeted anti
12	18	100.0	18	20 X23693	Deletion sequence

C	13	18	100.0	18	20 X27536	Synthetic RNA sequ
	14	18	100.0	18	20 X18702	Target bcl-2 antis
	15	18	100.0	18	20 V99434	Antisense oligonuc
	16	18	100.0	18	21 Z99003	CpG motif for immu
	17	18	100.0	18	21 Z60975	Nucleotide sequenc
	18	18	100.0	18	21 Z87997	BRE-labeled oligo
	19	18	100.0	18	21 Z98660	Human bcl-2 therap
	20	18	100.0	18	21 Z47643	Parasitic infectio
	21	18	100.0	18	21 Z47680	Parasitic infectio
	22	18	100.0	18	21 Z47850	Immunostimulatory
	23	18	100.0	18	21 Z47981	Immune remodeling
	24	18	100.0	18	21 Z48024	CpG-N motif SOS-OD
	25	18	100.0	20	20 V74246	Human Bcl-2 forwar
	26	18	100.0	27	17 T18388	Bcl-2 translation
	27	18	100.0	35	16 Q86644	Human bcl-2 oligon
	28	18	100.0	35	19 V19652	Human biallelic po
	29	18	100.0	251	19 X11646	Human biallelic po
	30	18	100.0	251	19 X12817	Human bcl-2 gene o
	31	18	100.0	615	16 Q73987	Human BCL2 cDNA
	32	18	100.0	760	17 T33694	Sequence of bcl-2
	33	18	100.0	765	14 Q49815	Bcl-2. Homo sapie
	34	18	100.0	831	9 N81293	Sequence of bcl-2
	35	18	100.0	911	20 X08431	bcl-2 proto-oncoge
	36	18	100.0	953	20 X33183	Bcl-2 DNA fragment
	37	18	100.0	5086	15 Q54631	Human oncogene bcl
	38	18	100.0	5086	16 Q86661	Human bcl-2 gene
	39	18	100.0	5086	19 X75766	Human bcl2 proto-o
	40	18	100.0	5105	9 N81292	Sequence of bcl-2
	41	18	100.0	7996	20 X33184	Base sequence of t
	42	17	94.4	17	19 V19659	Human bcl-2 antisense
	43	16.4	91.1	18	19 V52559	Unmethylated CpG d
	44	16.4	91.1	18	19 V47681	Unmethylated CpG d
	45	16.4	91.1	18	19 V27733	Immunostimulatory

ALIGNMENTS

RESULT	1
Q86659	ID Q86659 standard; DNA; 18 BP.
XX	Q86659;
AC	Q86659;
XX	27-SEP-1995 (first entry)
DT	Bcl-2 antisense oligonucleotide.
DE	Anticodon oligomer; antisense oligonucleotide; bcl-2; cancer; therapy;
XX	chemoresistance; ss.
KW	Synthetic.
XX	Key Location/Qualifiers
FT	misc_feature 1..18
FT	/*tag= a
FT	/note= "3'-5' (antisense) sequence"
XX	WO9508350-A.
PN	30-MAR-1995.
XX	20-SEP-1994; 94WO-US10725.
XX	20-SEP-1993; 93US-0124256.
PR	(REED/J) REED J C.
XX	Reed JC;
PA	WPI; 1995-139394/18.
XX	Anti-code oligomers which bind to bcl-2 mRNA - for the treatment
PT	

PT of human solid tumours, esp. breast cancer
 XX Example 18; Page 44; 108pp; English.
 PS
 XX Reversal of chemoresistance of tumor cells by antisense-mediated
 CC reduction of bcl1-2 expression was demonstrated using the
 CC oligonucleotide given in Q86659. This is antisense to the first
 CC 6 codons of the bcl-2 ORF.
 XX
 SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 16; Length 18;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
 |||||
 DB 1 tctcccagcgtgcgccat 18
 |||||

RESULT 2
 V52545
 ID V52545 standard; DNA; 18 BP.
 AC V52545;
 XX
 XX 20-NOV-1998 (first entry)
 DT
 XX
 DE Unmethylated CpG dinucleotide 1758.
 XX
 XX Unmethylated CpG dinucleotide; immune response; bacterial meningitis;
 KW natural killer cell activation; NK cell; Th2 response; neonatal sepsis;
 KW pulmonary disorder; asthma; environmentally induced airway disease;
 KW bacterial infection; endotoxaemia; therapy; cystic fibrosis;
 KW inflammatory bowel disease; ss.
 XX
 OS Synthetic.
 XX
 XX WO9837919-A1.
 PN
 XX
 PD 03-SEP-1998.
 XX
 XX 25-FEB-1998; 98WO-US03678.
 PF
 XX
 XX 28-FEB-1997; 97US-0039405.
 PR
 XX
 XX (IOWA) UNIV IOWA RES FOUND.
 PA
 XX Krieg AM, Schwartz DA;
 PI
 XX WPI; 1998-480941/41.
 DR
 XX
 XX Use of nucleic acids containing an unmethylated CpG - for treating a
 PT subject having or at risk of having an acute decrement in air flow
 PT or inhibiting an inflammatory response
 PT
 XX
 PS Example 4; Page 35; 65pp; English.
 XX
 XX This sequence represents an unmethylated CpG dinucleotide, and can be
 CC used in the method of the invention. The method is for treating a subject
 CC having, or at risk of having an acute decrement in air flow, comprising
 CC administering a nucleic acid sequence containing at least one
 CC unmethylated CpG. The nucleic acids containing an unmethylated CpG
 CC dinucleotide affect an immune response in a subject by activating natural
 CC killer cells (NK) or redirecting a subject's immune response from a Th2
 CC to a Th1 response by inducing monocytic and other cells to produce Th1
 CC cytokines. They can be used to treat pulmonary disorders having an
 CC immunologic component, such as asthma or environmentally induced airway
 CC disease. They can also be used to treat diseases associated with
 CC Gram-positive bacterial infections or endotoxaemia including bacterial
 CC meningitis, neonatal sepsis, cystic fibrosis, inflammatory bowel disease
 CC and liver cirrhosis, Gram-negative pneumonia, Gram-negative abdominal

CC abscess, haemorrhagic shock, disseminated intravascular coagulation, or
 CC an inflammatory response to lipopolysaccharide.
 XX
 SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
 |||||
 DB 1 tctcccagcgtgcgccat 18
 |||||

RESULT 3
 V27719
 ID V27719 standard; DNA; 18 BP.
 AC V27719;
 XX
 XX 01-OCT-1998 (first entry)
 DT
 XX
 DE Immunostimulatory oligodeoxyribonucleotide of the invention.
 XX
 XX Immunostimulatory; oligodeoxyribonucleotide; ODN;
 KW unmethylated CpG dinucleotide; activate; lymphocyte; immune response;
 KW Th2; Th1; cytokine; treatment; prevention; asthma; autoimmune disease;
 KW desensitisation therapy; artificial adjuvant; antibody generation; ss.
 XX
 OS Synthetic.
 XX
 XX WO9818810-A1.
 PN
 XX
 PD 07-MAY-1998.
 XX
 XX 30-OCT-1997; 97WO-US19791.
 PF
 XX
 XX 30-OCT-1996; 96US-0738652.
 PR
 XX
 XX (IOWA) UNIV IOWA RES FOUND.
 PA
 XX Kline JN, Krieg AM;
 PI
 XX WPI; 1998-272127/24.
 DR
 XX New immunostimulatory nucleic acid molecules - which contain at
 PT least one unmethylated CpG dinucleotide, used for treating e.g.
 PT tumours, infections or autoimmune disease
 PT
 XX
 PS Disclosure; Page 49; 109pp; English.
 XX
 XX V27641-751 represent immunostimulatory oligodeoxyribonucleotides (ODNs)
 CC of the invention. The ODNs contain at least one unmethylated CpG
 CC dinucleotide, and have the formula:
 CC 5' N1X1CX2N2 3', where at least one nucleotide separates consecutive
 CC CPGs, X1 is adenine, guanine or thymine, X2 is cytosine or thymine, N1
 CC does not contain a CCG tetramer or more than one CCG or CCG trimer OR
 CC any nucleotide and N1+N2 is 0-26 bases with the provision that N1 and N2
 CC are selected from GpT, GpG, GpA, Apt and Apa, X3 and X4
 CC are selected from Tpt or Cpt, N is any nucleotide and N1+N2 is 0-26
 CC bases with the provision that N1 and N2 does not contain a CCG tetramer
 CC or more than one CCG or CCG trimer.
 CC The ODNs activate lymphocytes in a subject and redirect a subject's
 CC immune response from a Th2 to a Th1 (e.g. by inducing monocytic cells and
 CC other cells to produce Th1 cytokines, including IL-12, IFN-gamma and
 CC GM-CSF). The ODNs can be used to treat or prevent an asthmatic disorder,
 CC autoimmune diseases, in desensitisation therapy, as an artificial
 CC adjuvant during antibody generation in a mammal such as a mouse or a
 CC human.
 XX
 XX Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | |
Db 1 tctcccagcgtgcgccat 18

RESULT 4

V28181
ID V28181 standard; DNA; 18 BP.

XX AC V28181;

XX DT 08-OCT-1998 (first entry)

XX DE Antisense oligonucleotide to bcl-2 mRNA.

XX KW Purification; oligonucleotide; matrix; affinity unit;

XX KW affinity purification; antisense; bcl-2; ss.

XX OS Synthetic.

XX PN WO9827425-A1.

XX PD 25-JUN-1998.

XX PF 18-DEC-1997; 97WO-US23284.

XX PR 19-DEC-1996; 96US-0769951.

XX PA (ISIS-) ISIS PHARM INC.

XX PI Chen D, Cole DL, Srivatsa GS;

XX DR WPI; 1998-362922/31.

XX PT Matrix for selective separation of oligo:nucleotide - useful for,
PT e.g. large scale purification of anti-sense agents from their
PT deletion derivatives formed during synthesis

XX PS Disclosure; Page 86; 183pp; English.

XX CC V28155-268 represent oligonucleotides which can be purified using the
CC method of the invention. The specification describes a matrix that
CC comprises a support and an affinity unit that specifically and reversibly
CC binds a target oligonucleotide, and comprises a sequence of bases having
CC the reverse complement of a hybridising portion of the target
CC oligonucleotide. The matrix is used for affinity purification of
CC synthetic oligonucleotides, specifically antisense agents, for treatment
CC of hyperproliferative diseases, for treating a non-pathogen,
CC non-hyperproliferative disease, e.g. Alzheimer's, for modulating
CC expression of cell surface proteins, and to inhibit a eukaryotic
CC pathogen, retrovirus or other viruses.

XX SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | |
Db 1 tctcccagcgtgcgccat 18

RESULT 5

V19667
ID V19667 standard; DNA; 18 BP.

XX V19667;

XX DT 12-JUN-1998 (first entry)

XX DE Human bcl-2 antisense oligonucleotide 13.

XX KW Antisense oligonucleotide; bcl-2 gene; lymphoma; leukaemia; human;
XX KW cancer; ss.

XX OS Synthetic.

XX OS Homo sapiens.

XX PN US5734033-A.

XX PD 31-MAR-1998.

XX PF 24-MAR-1994; 94US-0288692.

XX PR 21-FEB-1992; 92US-0840716.

XX PR 22-DEC-1988; 88US-0288692.

XX PR 24-MAR-1994; 94US-0217082.

XX PA (UYPE-) UNIV PENNSYLVANIA.

XX PI Reed J;

XX DR WPI; 1998-229881/20.

XX PT Anti-sense oligo:nucleotide(s) complementary to BCL-2 mRNA - useful
XX PT for treating cancers, e.g. lymphoma(s) and some leukaemia(s)

XX PS Disclosure; Column 23; 21pp; English.

XX CC This antisense oligonucleotide is complementary to the translation
XX CC initiation site of the human bcl-2 mRNA. The Bcl-2 antisense
XX CC oligonucleotides are phosphorothioate derivatives and can straddle
XX CC strategic sites such as the translation initiation site, donor and
XX CC acceptor splicing sites, or sites for transportation or degradation.
XX CC Blocking translation at such strategic sites prevents the formation of
XX CC a functional bcl-2 gene product. These oligonucleotides may be used for
XX CC treating cancers associated with high levels of bcl-2 gene expression,
XX CC especially lymphomas and some leukaemias.

XX SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 19; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | |
Db 1 tctcccagcgtgcgccat 18

RESULT 6

Z31944
ID Z31944 standard; DNA; 18 BP.

XX AC Z31944;

XX DT 26-JAN-2000 (first entry)

XX DE CpG adjuvant oligo 1002.

XX KW CpG adjuvant; vaccine; polyoxyethylene ether; polyoxyethylene ester;
XX KW antigen; infection; allergy; cancer; therapy; ss.

XX OS Synthetic.

XX PN WO9952549-A1.

XX XX

PD 21-OCT-1999.
 XX
 PF 29-MAR-1999; 99WO-EP02278.
 XX
 PR 09-APR-1998; 98GB-0007805.
 PR 25-SEP-1998; 98GB-0020956.
 XX
 PA (SMTK) SMITHKLINE BEECHAM BIOLOGICALS.
 XX
 PI Friede M, Hermand P;
 XX
 DR WPI; 1999-620290/53.
 XX
 PT Vaccine to protect against infections, allergy and cancer -
 XX
 XX Example 9; Page 26; 52pp; English.
 PS
 CC This sequence represents a CpG adjuvant that can be used in the vaccine
 CC composition of the invention. The vaccine comprises a polyoxyethylene
 CC ether or ester (I), not in the form of a vesicle, pharmaceutically
 CC acceptable excipient and an antigen (Ag) or antigenic composition. The
 CC vaccine can be used to treat or prevent infections (by bacteria, viruses
 CC or other parasites), allergy and cancer. (I), which are safe, easy to
 CC sterilize and simple to administer, are powerful vaccine adjuvants, able
 CC to induce a systemic immune response when administered (non-invasively)
 CC to the mucosa. The response is at least as good as that from conventional
 CC systemic injection. (I) are effective at low concentration, have low
 CC reactogenicity and are well tolerated.
 XX
 SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 tctcccagcgtgcgccat 18
 |||||
 Db 1 tctcccagcgtgcgccat 18

RESULT 7
 Z41905
 ID Z41905 standard; DNA; 18 BP.
 XX
 AC Z41905;
 XX
 DT 24-JAN-2000 (first entry)
 XX
 DE IL-12 secretion inducing CpG oligonucleotide 50.
 XX
 KW CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
 KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
 KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
 KW antigen presenting cell; infection; allergic disease.
 XX
 OS Synthetic.
 XX
 PN WO9951259-A2.
 XX
 PD 14-OCT-1999.
 XX
 PF 02-APR-1999; 99WO-US07335.
 XX
 PR 03-APR-1998; 98US-0080729.
 XX
 PA (IOWA) UNIV IOWA RES FOUND.
 XX
 PI Krieg AM, Weiner G;
 XX
 DR WPI; 1999-620169/53.
 XX
 PT Novel synergistic combinations of immunostimulatory oligonucleotides

PT and immunopotentiating cytokines are useful for stimulating the immune
 PT system -
 XX
 PS Example 8; Page 80; 91pp; English.
 XX
 CC Sequences Z41856-Z41949 are phosphorothioate CpG oligonucleotides which
 CC are used in the invention to induce interleukin-12 (IL-12) secretion
 CC from human PBMC. The invention comprises stimulating an immune response
 CC in a subject comprising administering to a subject exposed to an antigen,
 CC an immunopotentiating cytokine and an immunostimulatory CpG
 CC oligonucleotide to induce a synergistic antigen specific immune response.
 CC The methods are useful for treating cancer by stimulating an antigen
 CC specific immune response against a cancer antigen. The methods can also
 CC be used to treat neoplastic disorders in humans, including but not
 CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
 CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
 CC for treating infectious diseases, e.g. viral diseases such as HIV,
 CC bacterial diseases, and fungal diseases. The methods may also be used to
 CC treat allergic diseases, e.g. asthma. The methods and compositions may
 CC also be applied to treat cancer and tumours in non human subjects,
 CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
 CC be treated and include leukaemia, haemangiopericytoma and bovine ocular
 CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
 CC caused by the bacterium Corynebacterium pseudotuberculosis, and
 CC contagious lung tumour of sheep caused by jaagsiekte may also be treated.
 CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
 CC antigen presenting cells, such as monocytes and macrophages. CpG
 CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
 CC be used as an adjuvant in conjunction with tumour antigens to protect
 CC against a tumour challenge.
 XX

SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
 Best Local Similarity 100.0%; Pred. No. 1.5;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 tctcccagcgtgcgccat 18
 |||||
 Db 1 tctcccagcgtgcgccat 18

RESULT 8

Z41948
 ID Z41948 standard; DNA; 18 BP.

XX
 AC Z41948;

XX
 DT 24-JAN-2000 (first entry)

XX
 DE IL-12 secretion inducing CpG oligonucleotide 93.

XX
 DE CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
 KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
 KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
 KW antigen presenting cell; infection; allergic disease.

XX
 OS Synthetic.

XX
 PN WO9951259-A2.

XX
 PD 14-OCT-1999.

XX
 PF 02-APR-1999; 99WO-US07335.

XX
 PR 03-APR-1998; 98US-0080729.

XX
 PA (IOWA) UNIV IOWA RES FOUND.

XX
 PI Krieg AM, Weiner G;

XX
 DR WPI; 1999-620169/53.

XX Novel synergistic combinations of immunostimulatory oligonucleotides
PT and immunopotentiating cytokines are useful for stimulating the immune
PT system -
XX
XX Example 8; Page 88; 91pp; English.
XX
CC Sequences Z41856-Z41949 are phosphorothioate CpG oligonucleotides which
CC are used in the invention to induce interleukin-12 (IL-12) secretion
CC from human PBMC. The invention comprises stimulating an immune response
CC in a subject comprising administering to a subject exposed to an antigen,
CC an immunopotentiating cytokine and an immunostimulatory CpG
CC oligonucleotide to induce a synergistic antigen specific immune response.
CC The methods are useful for treating cancer by stimulating an antigen
CC specific immune response against a cancer antigen. The methods can also
CC be used to treat neoplastic disorders in humans, including but not
CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
CC for treating infectious diseases, e.g. viral diseases such as HIV,
CC bacterial diseases, and fungal diseases. The methods may also be used to
CC treat allergic diseases, e.g. asthma. The methods and compositions may
CC also be applied to treat cancer and tumours in non human subjects,
CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
CC be treated and include leukaemia, haemangiopericytoma and bovine ocular
CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
CC caused by the bacterium *Corynebacterium pseudotuberculosis*, and
CC contagious lung tumour of sheep caused by jaagsiekte may also be treated.
CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
CC antigen presenting cells, such as monocytes and macrophages. CpG
CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
CC be used as an adjuvant in conjunction with tumour antigens to protect
CC against a tumour challenge.
XX
XX Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 9
X78803
ID X78803 standard; DNA; 18 BP.
XX
AC X78803;
XX
DT 06-SEP-1999 (first entry)
XX
DE HPV fusion protein CpG oligonucleotide 2.

XX Fusion protein; E6 protein; E7 protein; E6/E7; immunomodulator; tumour;
KW immunological fusion partner; CpG oligonucleotide; immune response;
KW HPV antigen; prevention; treatment; primer; ss.
XX

OS Synthetic.
OS Human papillomavirus.

XX WO9933868-A2.

XX 08-JUL-1999.

XX 18-DEC-1998; 98WO-EP08563.

XX 24-DEC-1997; 97GB-0027262.

XX (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.

XX Dalemans WLJ, Gerard CMG;

XX WPI; 1999-405485/34.
XX
XX Composition comprising an E6, E7 or E6/E7 fusion protein from HPV to
PT induce immune response to HPV
XX
XX Claim 11; Page 37; 62pp; English.
XX
CC X78791-X78801 represent nucleic acid sequences which encode novel
CC constructs comprising an E6 or E7 protein or E6/E7 fusion protein from
CC HPV (represented in Y25375-Y25386). These constructs are optionally
CC linked to an immunological fusion partner and an immunomodulatory CpG
CC oligonucleotide. The products of the invention can be used to induce an
CC immune response in a patient to an HPV antigen. They can also be used
CC for preventing or treating HPV induced tumours. This sequence represents
CC a CpG oligonucleotide which is used in the method of the invention.
XX
XX Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 10
X88537
ID X88537 standard; DNA; 18 BP.

XX AC X88537;

XX 10-SEP-1999 (first entry)

DE Cytosine-guanosine dinucleotide motif oligonucleotide #4.

KW Cytosine-guanosine dinucleotide motif; CpG; immunomodulation;
KW unethylated; vaccine; immunostimulation; immune response;
KW T-independent type 1 antigen; T-independent type 2 antigen;
KW polysaccharide conjugate antigen; ss.

XX Synthetic.

XX WO9933488-A2.

XX 08-JUL-1999.

XX 18-DEC-1998; 98WO-EP08562.

XX 24-DEC-1997; 97GB-0027262.

XX (SMIK) SMITHKLINE BEECHAM BIOLOGICALS.

XX Dalemans WLJ, Laferriere CAJ, Prieels J;

XX WPI; 1999-405369/34.

XX A vaccine composition for inducing a immune response to
PT T-independent type 1 or type 2 antigen or polysaccharide conjugate
PT antigen
XX

XX Claim 6; Page 31; 35pp; English.

XX The present invention describes a formulation (A) comprising a
CC cytosine-guanosine dinucleotide motif (CpG) oligonucleotide and
CC T-independent type 1 or type 2 antigens or polysaccharide conjugate
CC antigen. The present sequence represent a specifically claimed CpG
CC oligonucleotide. A vaccine composition comprising the formulation is
CC used for inducing a immune response to T-independent type 1 or type 2
CC antigen or polysaccharide conjugate antigen. The use of

CC immunostimulatory CpG oligonucleotide acts as an adjuvant to
CC pneumococcal polysaccharides.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | |
Db 1 tctcccagcgtgcgccat 18

RESULT 11
X33514
ID X33514 standard; DNA; 18 BP.
XX
AC X33514;
XX
XX 07-JUL-1999 (first entry)
XX
DE BCL2-targeted antisense oligonucleotide SEQ ID NO:45.
XX
XX Combinatorial antisense library; oligonucleotide analogue; RNase;
KW ribozyme; cleavage; anchor; binding; target RNA; ss.
XX
OS Synthetic.
XX
XX WO9918238-A1.
XX
PN 15-APR-1999.
XX
XX 28-SEP-1998; 98WO-US20361.
XX
XX 18-AUG-1998; 98US-0136080.
PR
PR 02-OCT-1997; 97US-0060873.
XX
XX (OASIS-) OASIS BIOSCIENCES INC.
XX
XX Arnold LJ, Brown BD, Riley TA;
PI
XX WPI; 1999-264039/22.
DR

XX Oligonucleotide analog compositions capable of coupling to form
PT antisense molecules
PT
XX
PS Example 9; Page 45; 71pp; English.

XX The present invention describes a composition comprising two
CC oligonucleotide analogues, each having a binding domain and a coupling
CC moiety, where the binding domains are capable of hybridizing to a target
CC polynucleotide and the coupling moieties are capable of coupling to each
CC other in the absence of a target molecule. The composition/compound is
CC used to cleave an RNA target. The compositions can be used to determine
CC an optimal antisense site for a given mRNA or an optimal ribozyme
CC cleavage site for a target RNA. By separating the antisense molecules
CC into two or more pieces, a comprehensive antisense library can be
CC prepared in advance, rather than synthesizing a plurality of candidate
CC antisense molecules as needed. A complete library of every possible
CC 17-mer oligonucleotide, using the four natural bases, would consist of
CC 417 (or about 1.7 x 10¹⁰) molecules. By providing the antisense molecules
CC in at least two components, e.g. a library of 8-mers and a library of
CC 9-mers, assembled quickly as needed, the library size is reduced to 48 +
CC 49, or 327 650 molecules. The complexity of the library can be further
CC reduced by substituting one or more universal or degenerate bases for
CC some of the natural bases. The present sequence represents an
CC oligonucleotide, which is used in an example from the present invention.

XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | |
Db 1 tctcccagcgtgcgccat 18

RESULT 12
X23693
ID X23693 standard; DNA; 18 BP.
XX
AC X23693;
XX
XX 18-JUN-1999 (first entry)
XX
DE Deletion sequence oligonucleotide 146.
XX
XX
KW Deletion sequence oligonucleotide; sensor array; eukaryotic pathogen;
KW probe; cellular adhesion modulator; cellular proliferation modulator;
KW human retrovirus; human immunodeficiency virus; non-human retrovirus;
KW HIV; primer; ss.
XX
OS Synthetic.
XX
PN WO9911820-A1.
XX
PD 11-MAR-1999.
XX
XX 01-SEP-1998; 98WO-US18084.
PF
PR 02-SEP-1997; 97US-0923771.
XX
XX (ISTS-) ISIS PHARM INC.
XX
XX Chen D, Srivatsa GS;
PI
XX WPI; 1999-205198/17.
DR

XX New compositions comprising sensor arrays made up of unique probe
PT oligonucleotides - useful for characterizing a sample of target
PT deletion oligonucleotides
XX

PS Example 9; Page 152; 163pp; English.

XX This invention describes a novel composition comprising a number of
CC sensor arrays, where each array comprises a unique probe oligonucleotide,
CC which is the reverse complement of part of a unique target
CC oligonucleotide present in a mixture of target deletion sequence
CC oligonucleotides. The compositions form a method for characterizing a
CC sample of target deletion oligonucleotides which are labelled and
CC hybridize with the probe oligonucleotides of the sensor arrays. Such
CC oligonucleotides and their targets are represented in X23548-X23709.
CC Oligonucleotides characterized by the method form pharmaceutical
CC compositions that are useful for modulating cellular adhesion or
CC proliferation, and being active against a eukaryotic pathogen, a human
CC retrovirus, a human immunodeficiency virus (HIV), or a non-human
CC retrovirus, including influenza virus, Epstein-Barr virus, Respiratory
CC Syncytial Virus or cytomegalovirus (CMV). The compositions enable
CC characterization of deletion sequence oligonucleotides having related,
CC but different nucleobase sequences, and quantification of different
CC species of deletion sequence ("target") oligonucleotides in a mixture.
CC Also, if the specificity of the oligonucleotide's nucleobase sequence
CC for its reverse complement is not modified, the method may be performed
CC using oligodeoxynucleotides.

XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 13
X27536/c
ID X27536 standard; RNA; 18 BP.
XX
AC X27536;
XX
DT 27-MAY-1999 (first entry)
XX
DE Synthetic RNA sequence produced by the method of the invention.
XX
KW Silyloxymethyl; phosphonate; silyloxymethyl halide; diagnosis; ss;
KW cyanoethyl phosphoramidate coupling; isomerisation; steric hindrance.
XX
OS Synthetic.
XX
PN WO9909044-A1.
XX
PD 25-FEB-1999.
XX
PF 17-AUG-1998; 98WO-EP05215.
XX
PR 18-AUG-1997; 97CH-0001931.
XX
PA (JENNA/) JENNY L.
PA (PITS/) PITSCH S.
PA (WEIS/) WEISS P A.
XX
PI Jenny L, Pitsch S, Weiss PA;
XX
DR WPI; 1999-180963/15.
XX

PT 2-Silyloxymethyl ribonucleosides and their phosphonate derivatives
PT - have high purity, use in machine synthesis of ribonucleic acids,
PT enable longer oligonucleotide chain construction, and larger amounts
XX
PS Example 7; Page 26; 38pp; English.
XX
CC The invention relates to silyloxymethyl protected D- or L-ribonucleosides
CC and their phosphonates (I), and silyloxymethyl halides (II). (I) are
CC intermediates for synthesis of RNA-oligonucleotides with predetermined
CC nucleotide sequence, particularly by machine synthesis. The groups
CC specified above, apart from those on silyl, are those particularly for
CC the cyanoethyl phosphoramidate coupling. Uses of the oligoribonucleotide
CC products in diagnosis, therapy, and as research tools, are well known,
CC and are not dealt with in detail. (II) is an intermediate for (I). The
CC silyloxymethyl halide reagent is easy to prepare, and yields are high.
CC Introduction of the silyloxymethyl group into the ribonucleoside is
CC simple and rapid, and the acetal bond formed does not migrate,
CC eliminating particularly the prior art problem of 2' to 3' isomerisation.
CC The methylenedioxy group spacer between the silyl group and nucleoside
CC ring results in less steric hindrance than bulky direct silyloxy
CC linkages, enabling first, a range of choices for the silyl substituents,
CC to provide, e.g., acid or base stability; and second, higher yields in
CC coupling. Purer products are therefore obtained than in prior art,
CC enabling larger quantities and longer chains of oligoribonucleotides to
CC be synthesised successfully, and in shorter times.
XX
SQ Sequence 18 BP; 4 A; 4 C; 8 G; 2 U; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 tctcccagcgtgcgccat 18
|||||
Db 18 TCTCCCAGCGTGCGCCAT 1

RESULT 14
X18702
ID X18702 standard; DNA; 18 BP.
XX
AC X18702;
XX
DT 10-MAY-1999 (first entry)
XX
DE Target bcl-2 antisense oligonucleotide BCL-2.
XX
KW Cellular adhesion protein; proliferation; antisense oligonucleotide;
KW alimentary canal; transport; gastrointestinal mucosa; cancer;
KW Alzheimer's disease; beta-thalassemia; malaria; viral infection;
KW HIV; inflammation; ss.
XX
OS Synthetic.
XX
PN WO9901579-A1.
XX
PD 14-JAN-1999.
XX
PF 01-JUL-1998; 98WO-US13574.
XX
PR 01-JUL-1997; 97US-0886829.
XX
PA (ISIS-) ISIS PHARM INC.
XX
PI Hardee G, Teng C;
XX
DR WPI; 1999-106077/09.
XX
PT Composition comprising nucleic acid and penetration enhancer - used
PT particularly for delivering therapeutic antisense oligonucleotides
PT across the gastrointestinal mucosa, provides high bioavailability
XX
PS Example 2; Page 86; 115pp; English.
XX
CC A pharmaceutical composition has been developed which comprises a
CC nucleic acid and at least one penetration enhancer. The compositions are
CC used: (i) to treat or prevent any disease or disorder that can be
CC treated with the nucleic acid, e.g. cancer, Alzheimer's disease,
CC beta-thalassemia, malaria, viral infections (including human immune
CC deficiency virus (HIV)), inflammation, in human or animal medicine;
CC (ii) to investigate the role of a gene or gene product in non-human
CC animals; and (iii) to modulate gene expression in cells, tissues or
CC organs. The compositions provide bioavailability of at least 15,
CC preferably 17-35,%. The penetration enhancer improves: (i) transport of
CC the nucleic acid across the mucosa of the alimentary canal and into
CC cells; and (ii) increases stability of the nucleic acid. Oral
CC administration avoids the complications and expense of intravenous or
CC other methods of administration. X18669 to X18799 and X18801 represent
CC antisense oligonucleotides which can be used as the nucleic acid in
CC the method of the invention.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

RESULT 15
V99434
ID V99434 standard; DNA; 18 BP.
XX
AC V99434;

XX 22-MAR-1999 (first entry)
XX Antisense oligonucleotide directed against human bcl-2 gene.
XX
XX Antisense oligonucleotide; human bcl-2 gene; phosphorothioate;
KW phosphodiester; lipid-encapsulation; tumour; aberrant gene expression;
KW treatment; inflammation; infection; ss.
XX
XX Synthetic.
OS Homo sapiens.
XX
XX Key Location/Qualifiers
FT modified_base 1..18
FT /*tag= a
FT /note= "phosphorothioate or phosphodiester bonds"
XX
XX WO9851278-A2.
XX
XX 19-NOV-1998.
XX
XX 14-MAY-1998; 98WO-CA00485.
XX
XX 14-MAY-1997; 97US-0856374.
XX
XX (INEX-) INEX PHARM CORP.
XX
XX Ansell SM, Cullis P, Debeyer D, Harasym T, Hope MJ;
PI Klimuk SK, Scherrer P, Sempile SC;
XX
XX WPI; 1999-045179/04.
XX
XX Composition containing lipid-encapsulated therapeutic agent -
PT useful, e.g. for delivering antisense molecules or ribozymes or
PT treating diseases associated with aberrant gene expression
PT
XX
XX Disclosure; Page 23; 98pp; English.
XX
XX The present sequence represents an antisense oligonucleotide directed
CC against the human bcl-2 gene. The oligonucleotide can have either
CC phosphorothioate or phosphodiester bonds. The oligonucleotide is
CC lipid-encapsulated using the method of the invention. A composition
CC comprising lipid-encapsulated particles of a therapeutic agent,
CC e.g. antisense oligonucleotides, is prepared by mixing at least
CC 2 lipids with buffered aqueous solution of charged therapeutic
CC agent to form an intermediate mixture of lipid-encapsulated particles,
CC and changing the pH of the mixture to neutralise at least some of the
CC external surface charges on the particles. One lipid has a
CC (de)protonatable group with Ka such that the lipid is charged at a
CC first pH but neutral at a second pH (particularly near physiological pH)
CC and the buffer maintains this lipid in the charged form (i.e. cationic
CC when the therapeutic agent is anionic in the buffer, or vice versa). The
CC second lipid prevents particle aggregation during formation of the
CC lipid-therapeutic agent particles. The composition is used to introduce
CC therapeutic agents into cells, in vivo or in vitro, particularly to
CC treat or prevent diseases associated with aberrant gene expression in
CC mammals, specifically tumours, inflammation or infection.
XX
SQ Sequence 18 BP; 2 A; 8 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 18; DB 20; Length 18;
Best Local Similarity 100.0%; Pred. No. 1.5;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 tctcccagcgtgcgccat 18
|||||
Db 1 tctcccagcgtgcgccat 18

Search completed: December 4, 2000, 21:09:55
Job time: 16673 sec

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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 16:28:02 ; Search time 75.06 Seconds
(without alignments)
36.269 Million cell updates/sec

Title: US-09-369-941-1

Perfect score: 18

Sequence: 1 tctccagcgtgcgcacat 18

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 262060 seqs, 75620727 residues

Total number of hits satisfying chosen parameters: 524120

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued_Patents_NA:*

- 1: /cgn2.6/ptodata/2/ina/5A_COMB.seq:*
- 2: /cgn2.6/ptodata/2/ina/5B_COMB.seq:*
- 3: /cgn2.6/ptodata/2/ina/5C_COMB.seq:*
- 4: /cgn2.6/ptodata/2/ina/5D_COMB.seq:*
- 5: /cgn2.6/ptodata/2/ina/6_COMB.seq:*
- 6: /cgn2.6/ptodata/2/ina/PCRUS_COMB.seq:*
- 7: /cgn2.6/ptodata/2/ina/backfiles1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	18	2	US-08-217-082A-17
2	18	100.0	18	3	US-08-465-485A-17
3	18	100.0	18	3	US-08-465-485A-24
4	18	100.0	18	5	US-09-080-285-17
5	18	100.0	18	5	US-09-080-285-24
6	18	100.0	18	5	US-09-249-730-218
7	18	100.0	27	1	US-08-410-804-13
8	18	100.0	27	2	US-08-607-269-8
9	18	100.0	27	2	US-08-259-514-13
10	18	100.0	27	3	US-08-858-311-13
11	18	100.0	27	6	PCT-US95-04600-8
12	18	100.0	35	2	US-08-217-082A-2
13	18	100.0	35	3	US-08-465-485A-2
14	18	100.0	35	5	US-09-080-285-2
15	18	100.0	615	5	US-08-465-485A-22
16	18	100.0	615	5	US-09-080-285-22
17	18	100.0	623	7	5506344-3
18	18	100.0	717	3	US-08-465-485A-20
19	18	100.0	717	5	US-09-080-285-20
20	18	100.0	760	2	US-08-405-702A-11
21	18	100.0	831	7	5459251-3
22	18	100.0	831	7	5506344-4
23	18	100.0	911	6	PCT-US93-06251-3
24	18	100.0	1846	3	US-08-365-486A-16
25	18	100.0	4825	7	5459251-1
26	18	100.0	5086	3	US-08-465-485A-19

c 27	18	100.0	5086	3	US-08-365-486A-14	Sequence 14, Appl
c 28	18	100.0	5086	5	US-09-080-285-19	Sequence 19, Appl
c 29	18	100.0	5086	6	PCT-US93-05651-4	Sequence 4, Appl
c 30	18	100.0	5086	6	PCT-US93-06251-2	Sequence 2, Appl
c 31	18	100.0	5104	7	5506344-1	Patent No. 5506344
c 32	17	94.4	17	2	US-08-217-082A-9	Sequence 9, Appl
c 33	15.4	85.6	33	5	US-08-650-726-1	Sequence 1, Appl
c 34	15	83.3	17	2	US-08-217-082A-8	Sequence 8, Appl
c 35	15	83.3	17	4	US-08-877-831-1	Sequence 1, Appl
c 36	14.8	82.2	335	7	5175102-1	Patent No. 5175102
c 37	14.4	80.0	6002	2	US-08-698-551-15	Sequence 15, Appl
c 38	14.4	80.0	6002	3	US-08-602-228-15	Sequence 15, Appl
c 39	14.4	80.0	6002	3	US-08-839-032A-15	Sequence 15, Appl
c 40	14.4	80.0	34303	4	US-08-735-609-4	Sequence 4, Appl
c 41	14.4	80.0	34303	4	US-08-735-609-4	Sequence 4, Appl
c 42	14.4	80.0	34303	5	US-09-315-372-4	Sequence 4, Appl
c 43	14.4	80.0	34303	5	US-09-244-752-4	Sequence 4, Appl
c 44	14.4	80.0	34303	5	US-09-245-497-4	Sequence 4, Appl
c 45	14.4	80.0	34382	3	US-08-374-483-6	Sequence 6, Appl

ALIGNMENTS

RESULT 1
US-08-217-082A-17
; Sequence 17, Application US/08217082A
; Patent No. 5734033
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: ANTISENSE OLIGONUCLEOTIDES FOR INHIBITING THE
; GROWTH OF CELLS EXPRESSING THE HUMAN BCL-2 GENE
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; STREET: 224 Airport Parkway
; CITY: San Jose
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 95110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/217,082A
; FILING DATE: 24-MAR-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-067-55 FWC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: Synthetic DNA
US-08-217-082A-17

Query Match 100.0%; Score 18; DB 2; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 2
US-08-465-485A-17
; Sequence 17, Application US/08465485A
; Patent No. 5831066
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-465-485A-17

Query Match 100.0%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 3
US-08-465-485A-24
; Sequence 24, Application US/08465485A
; Patent No. 5831066

; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; ADDRESSEE: P.C.
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 24:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: Other nucleic acid;
; DESCRIPTION: Synthetic DNA
; ANTI-SENSE: YES
; FEATURE:
; NAME/KEY: Modified_base
; LOCATION: 16..17
; OTHER INFORMATION: Last two internucleoside linkages are
; OTHER INFORMATION: phosphorothioates
US-08-465-485A-24

Query Match 100.0%; Score 18; DB 3; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
| | | | | | | | | | | | | | | | | |
Db 1 TCTCCAGCGTGCGCCAT 18

RESULT 4
US-09-080-285-17
; Sequence 17, Application US/09080285
; Patent No. 6040181
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,

ADDRESSEE: P.C.
STREET: 1755 S. Jefferson Davis Hwy., Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/080,285
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/465,485
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/124,256
FILING DATE: 20-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/840,716
FILING DATE: 21-FEB-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/080,285
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/465,485
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/124,256
FILING DATE: 20-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/840,716
FILING DATE: 21-FEB-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/288,692
FILING DATE: 22-DEC-1988
ATTORNEY/AGENT INFORMATION:
NAME: Fortney, Andrew D.
REGISTRATION NUMBER: 34,600
REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
TELECOMMUNICATION INFORMATION:
TELEPHONE: (408) 436-2070
TELEFAX: (408) 436-2075
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-080-285-17

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 TCTCCAGCGTGCCTCAT 18

RESULT 5
US-09-080-285-24
Sequence 24, Application US/09080285
Patent No. 6040181
GENERAL INFORMATION:
APPLICANT: Reed, John
TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
ADDRESSEE: P.C.
STREET: 1755 S. Jefferson Davis Hwy., Suite 400
CITY: Arlington
STATE: Virginia
COUNTRY: U.S.A.
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/080,285
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/465,485
FILING DATE: 05-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/124,256
FILING DATE: 20-SEP-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/840,716
FILING DATE: 21-FEB-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/288,692
FILING DATE: 22-DEC-1988
ATTORNEY/AGENT INFORMATION:
NAME: Fortney, Andrew D.
REGISTRATION NUMBER: 34,600
REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
TELECOMMUNICATION INFORMATION:
TELEPHONE: (408) 436-2070
TELEFAX: (408) 436-2075
INFORMATION FOR SEQ ID NO: 24:
SEQUENCE CHARACTERISTICS:
LENGTH: 18 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: Other nucleic acid;
DESCRIPTION: Synthetic DNA
ANTI-SENSE: YES
FEATURE:
NAME/KEY: Modified_base
LOCATION: 16..17
OTHER INFORMATION: Last two internucleoside linkages are
OTHER INFORMATION: phosphorothioates
US-09-080-285-24

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
Db 1 TCTCCAGCGTGCCTCAT 18

RESULT 6
US-09-249-730-218
Sequence 218, Application US/09249730
Patent No. 6121000
GENERAL INFORMATION:
APPLICANT: WRIGHT, Jim A.
APPLICANT: YOUNG, Aiping H.
TITLE OF INVENTION: Antitumor Antisense Sequences Directed Against R1 and
TITLE OF INVENTION: R2 Components of Ribonucleotide Reductase
FILE REFERENCE: 032396-040
CURRENT APPLICATION NUMBER: US/09/249,730
CURRENT FILING DATE: 1999-02-11
NUMBER OF SEQ ID NOS: 220
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 218
LENGTH: 18
TYPE: DNA
ORGANISM: Human
US-09-249-730-218

Query Match 100.0%; Score 18; DB 5; Length 18;
Best Local Similarity 100.0%; Pred. No. 0.85;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
Db 1 tctcccagcgtgcgccat 18

RESULT 7

US-08-410-804-13/c

; Sequence 13, Application US/08410804

; Patent No. 5632994

; GENERAL INFORMATION:

; APPLICANT: Reed, John C.

; APPLICANT: Sato, Takaaki

; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS

; NUMBER OF SEQUENCES: 22

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Cathryn Campbell

; STREET: 4370 La Jolla Village Drive. Ste 700

; CITY: San Diego

; STATE: California

; COUNTRY: United States

; ZIP: 92122

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/410,804

; FILING DATE: 27-MAR-1995

; CLASSIFICATION: 435

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/259,514

; FILING DATE: 14-JUN-1994

; ATTORNEY/AGENT INFORMATION:

; NAME: Campbell, Cathryn

; REGISTRATION NUMBER: 31,815

; REFERENCE/DOCKET NUMBER: P-LJ 1389

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (619) 535-9001

; TELEFAX: (619) 535-8949

; INFORMATION FOR SEQ ID NO: 13:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 27 base pairs

; TYPE: nucleic acid

; STRANDEDNESS: single

; TOPOLOGY: linear

; MOLECULE TYPE: cDNA

US-08-410-804-13

Query Match

; Best Local Similarity 100.0%; Score 18; DB 1; Length 27;

; Mismatches 0; Conservative 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18

Db 25 TCTCCAGCGTGCGCCAT 8

RESULT 8

US-08-607-269-8/c

; Sequence 8, Application US/08607269

; Patent No. 5702897

; GENERAL INFORMATION:

; APPLICANT: Reed, John C.

; APPLICANT: Sato, Takaaki

; TITLE OF INVENTION: Interaction of Proteins Involved in a

; TITLE OF INVENTION: Cell Death Pathway

; NUMBER OF SEQUENCES: 29

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Campbell and Flores

; STREET: 4370 La Jolla Village Drive, Suite 700

; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/607,269
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/226,876
; FILING DATE: 13-APR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn A.
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-LJ 9882
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-607-269-8

Query Match 100.0%; Score 18; DB 2; Length 27;

; Best Local Similarity 100.0%; Pred. No. 0.86;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18

Db 25 TCTCCAGCGTGCGCCAT 8

RESULT 9

US-08-259-514-13/c

; Sequence 13, Application US/08259514

; Patent No. 5747245

; GENERAL INFORMATION:

; APPLICANT: Reed, John C.

; APPLICANT: Sato, Takaaki

; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS

; NUMBER OF SEQUENCES: 22

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Cathryn Campbell

; STREET: 4370 La Jolla Village Drive. Ste 700

; CITY: San Diego

; STATE: California

; COUNTRY: United States

; ZIP: 92122

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS

; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/08/259,514

; FILING DATE: 14-JUN-1994

; CLASSIFICATION: 435

; ATTORNEY/AGENT INFORMATION:

; NAME: Campbell, Cathryn

; REGISTRATION NUMBER: 31,815

; REFERENCE/DOCKET NUMBER: P-LJ 9954

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (619) 535-9001

; TELEFAX: (619) 535-8949

;; INFORMATION FOR SEQ ID NO: 13:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 27 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: single
;; TOPOLOGY: linear
;; MOLECULE TYPE: cDNA
US-08-259-514-13

Query Match 100.0%; Score 18; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
DB 25 TCTCCAGCGTGCGCCAT 8

RESULT 10
US-08-858-311-13/c
; Sequence 13, Application US/08858311
; Patent No. 5876939
; GENERAL INFORMATION:
; APPLICANT: Reed, John C.
; TITLE OF INVENTION: FAS ASSOCIATED PROTEINS
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Cathryn Campbell
; STREET: 4370 La Jolla Village Drive, Ste 700
; CITY: San Diego
; STATE: California
; COUNTRY: United States
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/858,311
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/410,804
; FILING DATE: 27-MAR-1995
; APPLICATION NUMBER: US 08/259,514
; FILING DATE: 14-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Campbell, Cathryn
; REGISTRATION NUMBER: 31,815
; REFERENCE/DOCKET NUMBER: P-LJ 1389
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-858-311-13

Query Match 100.0%; Score 18; DB 3; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
DB 25 TCTCCAGCGTGCGCCAT 8

RESULT 11
PCT-US95-04600-8/c
; Sequence 8, Application PC/TUS9504600
; GENERAL INFORMATION:
; APPLICANT: LA JOLLA CANCER RESEARCH FOUNDATION
; TITLE OF INVENTION: Interaction of Proteins Involved in
; TITLE OF INVENTION: a Cell Death Pathway
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Campbell and Flores
; STREET: 4370 La Jolla Village Drive, Suite 700
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92122
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/04600
; FILING DATE: 12-APR-1995
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Imbra, Richard J.
; REGISTRATION NUMBER: 37,643
; REFERENCE/DOCKET NUMBER: FP-LJ 1361
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 535-9001
; TELEFAX: (619) 535-8949
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
PCT-US95-04600-8

Query Match 100.0%; Score 18; DB 6; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgccat 18
|||||
DB 25 TCTCCAGCGTGCGCCAT 8

RESULT 12
US-08-217-082A-2/c
; Sequence 2, Application US/08217082A
; Patent No. 5734033
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: ANTISENSE OLIGONUCLEOTIDES FOR INHIBITING THE
; TITLE OF INVENTION: GROWTH OF CELLS EXPRESSING THE HUMAN BCL-2 GENE
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCLELLAND, MAIER & NEUSTADT,
; STREET: 224 Airport Parkway
; CITY: San Jose
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 95110
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25

; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/217,082A
; FILING DATE: 24-MAR-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-067-55 FWC
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ANTI-SENSE: NO
; US-08-217-082A-2

Query Match 100.0%; Score 18; DB 2; Length 35;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgcacat 18
|||||
Db 35 TCTCCCACCGTGC GCCAT 18

RESULT 13
US-08-465-485A-2/c
; Sequence 2, Application US/08465485A
; Patent No. 5831066
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/465,485A
; FILING DATE: 05-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.

; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ANTI-SENSE: NO
; US-08-465-485A-2

Query Match 100.0%; Score 18; DB 3; Length 35;
Best Local Similarity 100.0%; Pred. No. 0.86;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tctcccagcgtgcgcacat 18
|||||
Db 35 TCTCCCACCGTGC GCCAT 18

RESULT 14
US-09-080-285-2/c
; Sequence 2, Application US/09080285
; Patent No. 6040181
; GENERAL INFORMATION:
; APPLICANT: Reed, John
; TITLE OF INVENTION: Regulation of bcl-2 Gene Expression
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: OBLON, SPIVAK, MCCLELLAND, MAIER & NEUSTADT,
; STREET: 1755 S. Jefferson Davis Hwy., Suite 400
; CITY: Arlington
; STATE: Virginia
; COUNTRY: U.S.A.
; ZIP: 22202
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/080,285
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/465,485
; FILING DATE: 05-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/124,256
; FILING DATE: 20-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/840,716
; FILING DATE: 21-FEB-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/288,692
; FILING DATE: 22-DEC-1988
; ATTORNEY/AGENT INFORMATION:
; NAME: Fortney, Andrew D.
; REGISTRATION NUMBER: 34,600
; REFERENCE/DOCKET NUMBER: 3335-070-55 CONT
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (408) 436-2070
; TELEFAX: (408) 436-2075
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 35 base pairs
; TYPE: nucleic acid

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:06:39 ; Search time 1141.84 Seconds
(without alignments)
108.295 Million cell updates/sec

Title: US-09-369-941-2
Perfect score: 20
Sequence: 1 tccatgacgttcctcgtcgtt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 7189864 seqs, 3091403243 residues
Total number of hits satisfying chosen parameters: 14379728

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_est3:*
4: gb_est4:*
5: gb_est5:*
6: gb_est6:*
7: gb_est7:*
8: gb_est8:*
9: gb_est9:*
10: gb_est10:*
11: gb_est11:*
12: gb_est12:*
13: gb_est13:*
14: gb_est14:*
15: gb_est15:*
16: gb_est16:*
17: gb_est17:*
18: gb_est18:*
19: gb_est19:*
20: gb_est20:*
21: gb_est21:*
22: gb_est22:*
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58: em_esthum16:*
59: em_esthum17:*
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61: em_esthum19:*
62: em_esthum20:*
63: em_estin1:*
64: em_estin2:*
65: em_estin3:*
66: em_estin4:*
67: em_estom:*
68: em_estov1:*
69: em_estov2:*
70: em_estp11:*
71: em_estp12:*
72: em_estp13:*
73: em_estp14:*
74: em_estp15:*
75: em_estro1:*
76: em_estro2:*
77: em_estro3:*
78: em_estro4:*
79: em_estro5:*
80: em_estro6:*
81: em_estro7:*
82: em_estro8:*
83: em_estro9:*
84: em_estro10:*
85: em_estro11:*
86: em_estro12:*
87: em_estro13:*
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96: gb_gss5:*
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101: gb_gss10:*
102: gb_gss11:*
103: gb_gss12:*
104: gb_gss13:*
105: gb_gss14:*
106: gb_gss15:*
107: gb_gss16:*
108: gb_gss17:*
109: gb_gss18:*
110: gb_gss19:*
111: gb_gss20:*
112: gb_gss21:*
113: gb_gss22:*
114: gb_gss23:*
115: gb_gss24:*
116: em_gss5:*

117: em_gss6:*
 118: em_gss7:*
 119: em_gss8:*
 120: em_gss9:*
 121: em_gss10:*
 122: em_gss11:*
 123: em_gss12:*
 124: em_gss13:*
 125: em_gss14:*
 126: em_gss15:*
 127: em_gss16:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16.8	84.0	464	AA171941	AA171941.zp24f01.s
2	16.8	84.0	346	AM065908	AM065908.687002G08
3	16.8	84.0	797	CNS02N06	AL205647 Tetraodon
4	16.8	84.0	821	CNS03G84	AL242653 Tetraodon
5	16.8	84.0	992	CNS04Q04	AL269221 Tetraodon
6	16.8	84.0	994	CNS0421L	AL271542 Tetraodon
7	16.4	82.0	824	CNS01XP2	AL172016 Tetraodon
8	16.4	82.0	1015	CNS05MA3	AL345108 Tetraodon
9	16	80.0	179	AV525865	AV525865 AV525865
10	15.8	79.0	254	BB342084	BB342084 BB342084
11	15.8	79.0	262	BB398655	BB398655 BB398655
12	15.8	79.0	392	AI077366	AI077366 oy87ell.x
13	15.8	79.0	408	AI086210	AI086210 ow90d05.s
14	15.8	79.0	430	AM063142	AM063142 TNO287 KR
15	15.8	79.0	509	AO478992	AO478992 RPT-11-2
16	15.8	79.0	717	CNS04SXC	AL305773 Tetraodon
17	15.8	79.0	934	AO573722	AO573722 nxb0084B
18	15.8	79.0	938	CNS03I44	AL245101 Tetraodon
19	15.8	79.0	992	CNS04Q70	AL302229 Tetraodon
20	15.4	77.0	219	CNS021650	AL014523 F.rubripe
21	15.4	77.0	279	AO651891	AO651891 Sheared D
22	15.4	77.0	302	CNS023000	AL015860 F.rubripe
23	15.4	77.0	311	BB234999	BB234999 BB234999
24	15.4	77.0	317	CNS021653	AL014526 F.rubripe
25	15.4	77.0	347	CNS023012	AL015871 F.rubripe
26	15.4	77.0	395	CNS021632	AL014505 F.rubripe
27	15.4	77.0	467	AI181400	AI181400 uc59c05.r
28	15.4	77.0	537	AI1812316	AI1812316 1063 pine
29	15.4	77.0	552	CNS021654	AL014527 F.rubripe
30	15.4	77.0	554	CNS021629	AL014502 F.rubripe
31	15.4	77.0	580	CNS023009	AL015869 F.rubripe
32	15.4	77.0	584	CNS021635	AL014508 F.rubripe
33	15.4	77.0	597	CNS021641	AL014514 F.rubripe
34	15.4	77.0	598	CNS021637	AL014510 F.rubripe
35	15.4	77.0	609	CNS023992	AL014520 F.rubripe
36	15.4	77.0	612	CNS021647	AL014520 F.rubripe
37	15.4	77.0	629	BE195266	BE195266 HYSMER008
38	15.4	77.0	658	AI1234463	AI1234463 EST366 Ma
39	15.2	76.0	151	AA494709	AA494709 fa10h06.x
40	15.2	76.0	189	AI1771817	AI1771817 EST232917
41	15.2	76.0	253	BB385734	BB385734 BB385734
42	15.2	76.0	272	AA958394	AA958394 ual1c12.r
43	15.2	76.0	273	AO105312	AO105312 HS.3020_B
44	15.2	76.0	277	BB110967	BB110967 BB110967
45	15.2	76.0	278	BB230155	BB230155 BB230155

ALIGNMENTS

RESULT 1
 AA171941

LOCUS	AA171941	464 bp	mRNA	EST	23-DEC-1996
DEFINITION	zp24f01.s1	Stratagene neuroepithelium (#937231)	Homo sapiens cDNA		
Clone IMAGE:	610393	3'	similar to contains Alu repetitive element;		
mRNA sequence.					
ACCESSION	AA171941				
VERSION	AA171941.1	GI:1751000			
KEYWORDS					
SOURCE	human.				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiappelli,B., Chisoe,S., Dietrich,N., Dubuque,T., Favellio,A., Gish,W., Hawkins,M., Hultman,M., Kucaba,T., Lacy,M., Le,N., Maridis,E., Moore,B., Morris,M., Parsons,J., Prange,C., Rifkin,L., Roeding,T., Schellenberg,K., Soares,M.B., Tan,F., Thierry-Meg,J., Trevasakis,E., Underwood,K., Wohlmann,P., Waterston,R., Wilson,R. and Marra,M.				
TITLE	Generation and analysis of 280,000 human expressed sequence tags				
JOURNAL	Genome Res. 6 (9), 807-828 (1996)				
MEDLINE	97044478				
COMMENT	Contact: Wilson RK Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information. Seq primer: -40M13 fwd. from Amersham High quality sequence stop: 360.				
FEATURES	source				
location/Qualifiers	1..464				
/organism="Homo sapiens"					
/db_xref="GDB:4625652"					
/db_xref="taxon:9606"					
/clone="IMAGE:610393"					
/clone_lib="Stratagene neuroepithelium (#937231)"					
/dev_stage="Ntera-2/RA neuroepithelial cells"					
/lab_host="SOLR (kanamycin resistant)"					
/note="Vector: pBluescript SK-; site:1; EcoRI; Site:2; XhoI: Cloned unidirectionally. Primer: Oligo dT, NT2 cells (Ntera-2/cl.D1) induced with Retinoic Acid for 24 hours. Average insert size: 1.5 kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5' GAATTCGGCAGG 3' -3' adaptor sequence: 5' CTCGATTTTCTTCTTCTTCTT 3' "					
BASE COUNT	109 a 105 c 100 g 148 t				
ORIGIN					
Query Match	84.0%; Score 16.8; DB 2; Length 464;				
Best Local Similarity	90.0%; Pred. No. 2.3e+02;				
Matches	18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;				
Oy	1 tccatgacgtctcgtacgtt 20				
Db	431 tccatgacgttctcgtacgtt 450				
RESULT 2					
LOCUS	AM065908	546 bp	mRNA	EST	30-MAR-2000
DEFINITION	687002G08.y1 687 - Early embryo from Delaware Zea mays cDNA, mRNA				
ACCESSION	AM065908				
VERSION	AM065908.1	GI:6020980			
KEYWORDS					
SOURCE	Zea mays.				
ORGANISM	Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Zea.				
REFERENCE	1 (bases 1 to 546)				
AUTHORS	Walbot,V.				

TITLE Maize ESTs from various cDNA libraries sequenced at Stanford University
JOURNAL Unpublished (1999)
COMMENT Contact: Walbot V
Department of Biological Sciences
Stanford University
855 California Ave, Palo Alto, CA 94304, USA
Tel: 650 723 8227
Fax: 650 723 8221
Email: walbot@stanford.edu
Plate: 687002 row: G column: 08.
FEATURES
source
1..546
/organism="Zea mays"
/cultivar="Illinois High Oil"
/db_xref="taxon:4577"
/clone_1lb="687 - Early embryo from Delaware"
/tissue_type="embryo"
/dev_stage="14, 21, 28, and 35 days after pollination"
/lab_host="E. coli SOLR"
/note="Organ: embryo; Vector: pBluescript SK; Site: 1: XhoI ; Site: 2: EcoRI; Library was prepared by Stratagene using the Uni-ZAP XR system (Stratagene BN937328-12). Clones were picked by a Q-bot after blue/white selection (ampicillin resistance - use 100 micrograms/microliter). Developed from a pool of equal amounts of RNA from developing embryos sampled at 14, 21, 28 and 35 days after pollination of the Illinois High Oil Maize Strain Cycle 90. This closed strain has been selected for high oil concentration for 90 generations and originates from the 1890s era open pollinated variety Burr's White"
113 a 183 c 156 g 94 t
BASE COUNT
ORIGIN
Query Match 84.0%; Score 16.8; DB 19; Length 546;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
RESULT 3
LOCUS CNS02N06/c 797 bp DNA GSS 14-MAY-2000
DEFINITION Tetradon nigroviridis genome survey sequence T7 end of clone 151108 of library G from Tetradon nigroviridis, genomic survey sequence.
ACCESSION AL205647.1 GI:7864466
VERSION GSS; genome survey sequence.
KEYWORDS Tetradon nigroviridis.
SOURCE Tetradon nigroviridis
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Eurypterygii; Ctenosquamata; Acanthomorpha; Eucanthomorpha; Holacanthopterygii; Acanthopterygii; Percomorpha; Tetraodontiformes; Tetraodontidae; Tetraodon.
REFERENCE 1 (bases 1 to 797)
Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 797)
Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Wincker,P., Brotlier,P., Quetier,F., Saurin,W. and Weissenbach,J.
AUTHORS Human gene number estimate provided by genome wide analysis using Tetradon nigroviridis DNA sequence

JOURNAL Unpublished
REFERENCE 3 (bases 1 to 797)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetradon nigroviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tetradon>.
FEATURES
source
1..797
/organism="Tetradon nigroviridis"
/db_xref="taxon:99883"
/clone_1lb="151108"
/clone_1lb="G"
/note="Genoscope sequence ID : CGAG151DF04LP1-end : T7"
BASE COUNT 192 a 204 c 209 g 177 t 15 others
ORIGIN
Query Match 84.0%; Score 16.8; DB 113; Length 797;
Best Local Similarity 90.0%; Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
RESULT 4
LOCUS CNS03G84 821 bp DNA GSS 17-MAY-2000
DEFINITION Tetradon nigroviridis genome survey sequence T7 end of clone 023N02 of library G from Tetradon nigroviridis, genomic survey sequence.
ACCESSION AL242653.1 GI:7963422
VERSION GSS; genome survey sequence.
KEYWORDS Tetradon nigroviridis.
SOURCE Tetradon nigroviridis
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Holacanthopterygii; Acanthomorpha; Eucanthomorpha; Tetraodontiformes; Tetraodontidae; Tetraodon.
REFERENCE 1 (bases 1 to 821)
Roest-Crollius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C., Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetradon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 821)
Roest-Crollius,H., Jalllon,O., Dasilva,C., Bouneau,L., Fisher,C., Bernot,A., Fizames,C., Wincker,P., Brotlier,P., Quetier,F., Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using Tetradon nigroviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 821)
Genoscope.
AUTHORS Direct Submission
TITLE Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
COMMENT This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetradon nigroviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tetradon>.
FEATURES
source
1..821
/organism="Tetradon nigroviridis"
/db_xref="taxon:99883"
/clone_1lb="023N02"
/clone_1lb="G"

BASE COUNT 217 a 145 c 172 g 279 t 8 others
ORIGIN

Query Match 84.0%: Score 16.8; DB 114; Length 821;
Best Local Similarity 90.0%: Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
||| ||||| ||||| |||||
Db 627 tccctgacgttactgacgtt 646

RESULT 5
CNS04004 992 bp DNA GSS 18-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 073A13 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL269221.1 GI:7991098
VERSION AL269221.1
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthopterygii; Percormorpha;
Holacanthopterygii; Acanthopterygii; Tetraodontiformes; Tetraodon.
1 (bases 1 to 992)
Roest-Crolius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bonneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Unpublished
2 (bases 1 to 992)
Roest-Crolius,H., Jalllon,O., Dasilva,C., Bonneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
Unpublished
3 (bases 1 to 992)
Genoscope.
REFERENCE Direct Submission
AUTHORS Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
TITLE This sequence is a single read and was generated as part of a large
JOURNAL scale clone-end sequencing project of the Tetraodon nigroviridis
COMMENT genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
Location/Qualifiers
FEATURES
source
1..992
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/db_xref="taxon:99883"
/clone_1bp="G"
/clone_11bp="G"
/note="Genoscope sequence ID : COBG073AA07LPI-end : T7"

BASE COUNT 242 a 184 c 211 g 337 t 18 others
ORIGIN

Query Match 84.0%: Score 16.8; DB 114; Length 992;
Best Local Similarity 90.0%: Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
||| ||||| ||||| |||||
Db 815 tccctgacgttactgacgtt 834

RESULT 6

CNS0421L 994 bp DNA GSS 18-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 077D02 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL271542.1 GI:7993521
VERSION AL271542.1
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthopterygii; Percormorpha;
Holacanthopterygii; Acanthopterygii; Tetraodontiformes; Tetraodon.
1 (bases 1 to 994)
Roest-Crolius,H., Jalllon,O., Dasilva,C., Fizames,C., Fisher,C.,
Bonneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Unpublished
2 (bases 1 to 994)
Roest-Crolius,H., Jalllon,O., Dasilva,C., Bonneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
Unpublished
3 (bases 1 to 994)
Genoscope.
REFERENCE Direct Submission
AUTHORS Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
TITLE This sequence is a single read and was generated as part of a large
JOURNAL scale clone-end sequencing project of the Tetraodon nigroviridis
COMMENT genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
Location/Qualifiers
FEATURES
source
1..994
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/db_xref="taxon:99883"
/clone="077D02"
/clone_1bp="G"
/note="Genoscope sequence ID : COBG077B01LPI-end : T7"

BASE COUNT 309 a 199 c 226 g 256 t 4 others
ORIGIN

Query Match 84.0%: Score 16.8; DB 114; Length 994;
Best Local Similarity 90.0%: Pred. No. 2.5e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 tccatgacgttcctgacgtt 20
||| ||||| ||||| |||||
Db 206 tccatgacgttccagccgtt 225

RESULT 7
CNS01XP2 824 bp DNA GSS 12-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence PUC-ori end of clone
DEFINITION 202P06 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL172016.1 GI:7810073
VERSION AL172016.1
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthopterygii; Percormorpha;
Holacanthopterygii; Acanthopterygii; Tetraodontiformes; Tetraodon.

REFERENCE 1 (bases 1 to 824)
AUTHORS Roest-Crolius,H., Jallion,O., Dasilva,C., Fizames,C., Fisher,C.,
Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 824)
AUTHORS Roest-Crolius,H., Jallion,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 824)
AUTHORS Genoscope.
TITLE Direct Submission
COMMENT Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetraodon>.
location/Qualifiers
1. 824
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="202P06"
/clone_1lb="g"
/note="Genoscope sequence ID : C0AG202DH03SP1-end :
PVC-0r1"

BASE COUNT 167 a 236 c 287 g 104 t 30 others
ORIGIN

Query Match 82.0%; Score 16.4; DB 113; Length 824;
Best Local Similarity 94.4%; Pred. No. 3.9e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 tccatgacgttcctgacg 18
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Db 232 tccatcacgttcctgcgacg 249

RESULT 8
CNS05N3A3 1015 bp DNA GSS 26-MAY-2000
LOCUS Tetraodon nigroviridis genome survey sequence SP6 end of clone
DEFINITION 035P01 of library B from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL345108
VERSION AL345108.1 GI:8238878
KEYWORDS GSS: genome survey sequence.
SOURCE Tetraodon nigroviridis.
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Eurypterygii; Ctenosquamata; Acanthomorphi; Euacanthomorphi;
Holacanthopterygii; Acanthopterygii; Percomorpha;
Tetraodontiformes; Tetraodontidae; Tetraodontidae; Tetraodon.
1 (bases 1 to 1015)
Roest-Crolius,H., Jallion,O., Dasilva,C., Fizames,C., Fisher,C.,
Bouneau,L., Billault,A., Quetier,F., Saurin,W., Bernot,A. and
Weissenbach,J.
TITLE Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1015)
AUTHORS Roest-Crolius,H., Jallion,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,W. and Weissenbach,J.
TITLE Human gene number estimate provided by genome wide analysis using
Tetraodon nigroviridis DNA sequence
JOURNAL Unpublished

REFERENCE 3 (bases 1 to 1015)
AUTHORS Genoscope.
TITLE Direct Submission
COMMENT Submitted (12-APR-2000) to the EMBL/GenBank/DBJ databases
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
<http://www.genoscope.cns.fr/Tetraodon>.
location/Qualifiers
1. 1015
/organism="Tetraodon nigroviridis"
/db_xref="taxon:99883"
/clone="035P01"
/clone_1lb="g"
/note="Genoscope sequence ID : C0RB035CH01B1-end : SP6"

BASE COUNT 253 a 268 c 261 g 214 t 19 others
ORIGIN

Query Match 82.0%; Score 16.4; DB 115; Length 1015;
Best Local Similarity 94.4%; Pred. No. 4e+02;
Matches 17; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 3 catgacgttcctgacgt 20
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Db 234 catgacgttcctgacgt 217

RESULT 9
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LOCUS AV525865 Arabidopsis thaliana aboveground organs two to six-week
DEFINITION Old Arabidopsis thaliana cDNA clone APD31f02R 5', mRNA sequence.
ACCESSION AV525865
VERSION AV525865.1 GI:8685393
KEYWORDS EST.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Embryophyta; Tracheophyta; Spermatophyta;
Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II;
Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 179)
Asamizu,E., Nakamura,Y., Sato,S. and Tabata,S.
A large scale analysis of cDNA in Arabidopsis thaliana: Generation
of 12,028 non-redundant expressed sequence tags from normalized and
size-selected cDNA libraries
DNA Res. 7, 175-180 (2000)
Contact: Erika Asamizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asamizu@kazusa.or.jp, URL:<http://www.kazusa.or.jp/en/plant/>.
location/Qualifiers
1. 179
/organism="Arabidopsis thaliana"
/strain="Columbia"
/db_xref="taxon:3702"
/clone="APD31f02R"
/clone_1lb="Arabidopsis thaliana aboveground organs two to
six-week old"
/tissue_type="aboveground organs"
/dev_stage="two to six-week old"
/note="Vector: phuescriptptII SK-, Site_1: EcoRI; Site_2:
XhoI"

BASE COUNT 47 a 35 c 48 g 49 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 5.3e+02;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 5 tgacgttcctgacgt 20

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:08:04 ; Search time 75.06 Seconds
(without alignments)
40.299 Million cell updates/sec

Title: US-09-369-941-2

Perfect score: 20

Sequence: 1 tccatgacgtctcgtcgtt 20

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 262060 seqs, 75620727 residues

Total number of hits satisfying chosen parameters: 524120

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued_Patents_NA:*

- 1: /cgn2_6/ptodata/2/ina/5A_COMB.seq:*
- 2: /cgn2_6/ptodata/2/ina/5B_COMB.seq:*
- 3: /cgn2_6/ptodata/2/ina/5C_COMB.seq:*
- 4: /cgn2_6/ptodata/2/ina/5D_COMB.seq:*
- 5: /cgn2_6/ptodata/2/ina/6_COMB.seq:*
- 6: /cgn2_6/ptodata/2/ina/PCtUS_COMB.seq:*
- 7: /cgn2_6/ptodata/2/ina/bcKilled1.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	4	US-09-133-774-12
2	20	100.0	20	5	US-09-303-862-12
3	16.8	84.0	20	4	US-09-133-774-11
4	16.8	84.0	20	5	US-08-386-063-25
5	16.8	84.0	20	5	US-09-303-862-11
6	16.8	84.0	2470	1	US-07-745-206A-14
7	16.8	84.0	2470	3	US-08-311-363-14
8	16.8	84.0	5467	1	US-07-745-206A-12
9	16.8	84.0	5467	1	US-08-311-363-12
10	16.8	84.0	7175	2	US-08-455-543A-8
11	16.8	84.0	7175	2	US-08-193-078B-8
12	16.8	84.0	7175	3	US-08-223-305C-8
13	16.8	84.0	7175	3	US-08-149-097D-8
14	16.8	84.0	7175	3	US-08-949-386-8
15	16.8	84.0	7175	5	US-08-450-562-8
16	16.8	84.0	7266	5	US-08-713-118-1
17	16.8	84.0	7362	2	US-08-455-543A-7
18	16.8	84.0	7362	2	US-08-193-078B-7
19	16.8	84.0	7362	3	US-08-223-305C-7
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21	16.8	84.0	7362	5	US-08-949-386-7
22	16.8	84.0	7362	5	US-08-450-562-7
23	15.8	79.0	8931	5	US-09-028-934-28
24	15.2	76.0	1799	3	US-08-560-398-3
25	14.8	74.0	840	1	US-07-906-983-1
26	14.4	72.0	700	5	US-08-764-563-2

27	14.2	71.0	29	2	US-08-484-557C-8	Sequence 8, Appl
28	14.2	71.0	29	2	US-08-487-426B-8	Sequence 8, Appl
29	14.2	71.0	29	3	US-08-487-720A-8	Sequence 8, Appl
30	14.2	71.0	76	6	PCT-US96-09451-8	Sequence 8, Appl
31	14.2	71.0	77	5	US-08-945-734-8	Sequence 8, Appl
32	14.2	71.0	355	7	5244792-11	Patent No. 5244792
33	14.2	71.0	523	3	US-08-628-413-1	Sequence 1, Appl
34	14.2	71.0	1949	2	US-08-760-335A-1	Sequence 1, Appl
35	14.2	71.0	1979	2	US-08-392-828C-3	Sequence 3, Appl
36	14.2	71.0	1979	5	US-09-330-945-3	Sequence 3, Appl
37	14.2	71.0	2019	3	US-08-455-073A-5	Sequence 5, Appl
38	14.2	71.0	8140	1	US-08-297-294A-1	Sequence 1, Appl
39	14.2	71.0	50341	2	US-08-247-901C-1	Sequence 1, Appl
40	14.2	71.0	50341	4	US-09-075-904-1	Sequence 1, Appl
41	13.8	69.0	15239	2	US-08-390-878-17	Sequence 17, Appl
42	13.6	68.0	20	1	US-08-436-714-7	Sequence 7, Appl
43	13.6	68.0	20	1	US-08-442-705-7	Sequence 7, Appl
44	13.6	68.0	20	2	US-08-337-829-7	Sequence 7, Appl
45	13.6	68.0	20	5	US-08-386-063-21	Sequence 21, Appl

ALIGNMENTS

RESULT 1

US-09-133-774-12

Sequence 12, Application US/09133774B

Patent No. 5962636

GENERAL INFORMATION:

APPLICANT: Bachmaier, Kurt

APPLICANT: Hessel, Andrew J.

APPLICANT: Neu M.D., Nikolaus

APPLICANT: Penninger, Josef M.

TITLE OF INVENTION: No. 5962636el Peptides Capable of Modulating Inflammatory Hear

FILE REFERENCE: A-536

CURRENT APPLICATION NUMBER: US/09/133,774B

CURRENT FILING DATE: 1998-08-12

NUMBER OF SEQ ID NOS: 26

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 12

LENGTH: 20

TYPE: DNA

ORGANISM: Chlamydia trachomatis

FEATURE:

OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a

OTHER INFORMATION: 60 kda cysteine rich outer membrane protein from

US-09-133-774-12

Query Match 100.0%; Score 20; DB 4; Length 20;

Best local Similarity 100.0%; Pred. No. 0.089;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgacgtctcgtcgtt 20

Db 1 tccatgacgtctcgtcgtt 20

RESULT 2

US-09-303-862-12

Sequence 12, Application US/09303862

Patent No. 6034230

GENERAL INFORMATION:

APPLICANT: Bachmaier, Kurt

APPLICANT: Hessel, Andrew J.

APPLICANT: Neu M.D., Nikolaus

APPLICANT: Penninger, Josef M.

TITLE OF INVENTION: No. 6034230el Peptides Capable of Modulating Inflammatory Hear

FILE REFERENCE: A-536

CURRENT APPLICATION NUMBER: US/09/303,862

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; CURRENT FILING DATE: 1999-05-03
; EARLIER APPLICATION NUMBER: 09/133,774
; EARLIER FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 12
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-303-862-12
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Query Match          100.0%; Score 20; DB 5; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.089;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 tccatgacgttcctcgaagctt 20
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RESULT 3
US-09-133-774-11
; Sequence 11, Application US/09133774B
; Patent No. 5962636
; GENERAL INFORMATION:
; APPLICANT: Bachmaler, Kurt
; APPLICANT: Hessel, Andrew J.
; APPLICANT: Neu M.D., Nikolaus
; APPLICANT: Penninger, Josef M.
; TITLE OF INVENTION: No. 5962636el Peptides Capable of Modulating Inflammatory Heart
; FILE REFERENCE: A-536
; CURRENT APPLICATION NUMBER: US/09/133,774B
; CURRENT FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 11
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-133-774-11
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Query Match          84.0%; Score 16.8; DB 4; Length 20;
Best Local Similarity 90.0%; Pred. No. 3.5;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 tccatgacgttcctcgaagctt 20
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RESULT 4
US-08-386-063-25
; Sequence 25, Application US/08386063
; Patent No. 6008200
; GENERAL INFORMATION:
; APPLICANT: Arthur M. Kriegl, M.D.
; TITLE OF INVENTION: IMMUNOMODULATORY OLIGONUCLEOTIDES
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LAHIVE & COCKFIELD
; STREET: 60 STATE STREET, SUITE 510
; CITY: BOSTON
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; STATE: MASSACHUSETTS
; COUNTRY: USA
; ZIP: 02109-1875
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/386,063
; FILING DATE:
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: ARNOLD, BETH E.
; REGISTRATION NUMBER: 35,430
; REFERENCE/DOCKET NUMBER: UI2-013CP
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)227-7400
; TELEFAX: (617)227-5941
; INFORMATION FOR SEQ ID NO: 25:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
US-08-386-063-25
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Query Match          84.0%; Score 16.8; DB 5; Length 20;
Best Local Similarity 90.0%; Pred. No. 3.5;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 tccatgacgttcctcgaagctt 20
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RESULT 5
US-09-303-862-11
; Sequence 11, Application US/09303862
; Patent No. 6034230
; GENERAL INFORMATION:
; APPLICANT: Bachmaler, Kurt
; APPLICANT: Hessel, Andrew J.
; APPLICANT: Neu M.D., Nikolaus
; APPLICANT: Penninger, Josef M.
; TITLE OF INVENTION: No. 6034230el Peptides Capable of Modulating Inflammatory Hear
; FILE REFERENCE: A-536
; CURRENT APPLICATION NUMBER: US/09/303,862
; CURRENT FILING DATE: 1999-05-03
; EARLIER APPLICATION NUMBER: 09/133,774
; EARLIER FILING DATE: 1998-08-12
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 11
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia trachomatis
; FEATURE:
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from
; OTHER INFORMATION: Chlamydia trachomatis.
US-09-303-862-11
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Query Match          84.0%; Score 16.8; DB 5; Length 20;
Best Local Similarity 90.0%; Pred. No. 3.5;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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Db 1 tccatgacgttcctgatgct 20

RESULT 6
US-07-745-206A-14/c
; Sequence 14, Application US/07745206A
; Patent No. 5429921
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Feldman, Daniel
; TITLE OF INVENTION: Human Calcium Channel Compositions and
; TITLE OF INVENTION: Methods
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fitch, Even, Tabin & Flannery
; STREET: 135 S. LaSalle
; CITY: Chicago
; STATE: Illinois
; COUNTRY: U.S.A.
; ZIP: 60603
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/745,206A
; FILING DATE: 19910815
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Feder, Scott B
; REFERENCE/DOCKET NUMBER: 51504
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 312-372-7842
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2470 base pairs
; TYPE: NUCLEIC ACID
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..2469
; US-07-745-206A-14

Query Match 84.0%; Score 16.8; DB 1; Length 2470;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgaagtt 20
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Db 509 tccatgacgttcacgccgtt 490

RESULT 7
US-08-311-363-14/c
; Sequence 14, Application US/08311363
; Patent No. 5876958
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: Feldman, Daniel
; APPLICANT: McCue, Ann
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: Human Calcium Channel Compositions and
; TITLE OF INVENTION: Methods
; NUMBER OF SEQUENCES: 32

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: USA
; ZIP: 92101-2926
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/311,363
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/745,206
; FILING DATE: 15-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 6362-51506
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619)238-0999
; TELEFAX: (619)238-0062
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2470 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..2469
; US-08-311-363-14

Query Match 84.0%; Score 16.8; DB 3; Length 2470;
Best Local Similarity 90.0%; Pred. No. 6.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgaagtt 20
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Db 509 tccatgacgttcacgccgtt 490

RESULT 8
US-07-745-206A-12/c
; Sequence 12, Application US/07745206A
; Patent No. 5429921
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Feldman, Daniel
; TITLE OF INVENTION: Human Calcium Channel Compositions and
; TITLE OF INVENTION: Methods
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fitch, Even, Tabin & Flannery
; STREET: 135 S. LaSalle
; CITY: Chicago
; STATE: Illinois
; COUNTRY: U.S.A.
; ZIP: 60603
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/07/745,206A
FILING DATE: 19910815
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Feder, Scott B
REFERENCE/DOCKET NUMBER: 51504
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312-372-7842
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 5467 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390..3392, 3396..3488, 3495..3539, 3543..3581, 3585..3587, 3591..3626, 3630..3689, 3693..3737, 3744..3746, 3750..4823, 4827..4841, 4845..5006, 5010..5096, 5100..5306, 5310..5366, 5370..5465)
US-07-745-206A-12

Query Match 84.0%; Score 16.8; DB 1; Length 5467;
Best Local Similarity 90.0%; Pred. No. 7.3;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgtcctgacgtt 20
|||||
Db 652 tccatgacgtcctgacgtt 633

RESULT 9
US-08-311-363-12/c
Sequence 12, Application US/08311363
Patent No. 5876958
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert
TITLE OF INVENTION: Human Calcium Channel Compositions and
TITLE OF INVENTION: Methods
NUMBER OF SEQUENCES: 32
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/311,363
FILING DATE:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: Seligman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-51506
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619)238-0999
TELEFAX: (619)238-0062

INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 5467 base pairs
TYPE: nucleic acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390..3392, 3396..3488, 3495..3539, 3543..3581, 3585..3587, 3591..3626, 3630..3689, 3693..3737, 3744..3746, 3750..4823, 4827..4841, 4845..5006, 5010..5096, 5100..5306, 5310..5366, 5370..5465).
US-08-311-363-12

Query Match 84.0%; Score 16.8; DB 3; Length 5467;
Best Local Similarity 90.0%; Pred. No. 7.3;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgtcctgacgtt 20
|||||
Db 652 tccatgacgtcctgacgtt 633

RESULT 10
US-08-455-543A-8/c
Sequence 8, Application US/0845543A
Patent No. 5792846
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert
TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
TITLE OF INVENTION: METHODS
NUMBER OF SEQUENCES: 57
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/455,543A
FILING DATE: May 31, 1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/223,305
FILING DATE: April 4, 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/868,354
FILING DATE: April 10, 1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/620,250
FILING DATE: 30-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/482,384
FILING DATE: 20-FEB-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/603,751
FILING DATE: 04-APR-1989

PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US89/01408
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/176,899
FILING DATE: 04-APR-1988
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-52517
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619)238-0062
TELEFAX: (619)238-0062
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 7175 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 144..6857
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..143
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 6855..7175
US-08-455-543A-8

Query Match 84.0%; Score 16.8; DB 2; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgagcttcctcagcgtt 20
|||||
DB 652 tccatgacgttcacgccggt 633

RESULT 11
US-08-193-078B-8/C
Sequence 8, Application US/08193078B
Patent No. 5846757
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROWN, MARTIN, HALLER & MCCLAIN
STREET: 1660 UNION STREET
CITY: SAN DIEGO
STATE: CA
COUNTRY: USA
ZIP: 92101
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentia Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/193,078B
FILING DATE: 07-FEB-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/868,354

FILING DATE: 10-APR-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-53607
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619-238-0062
TELEFAX: 619-238-0062
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 7175 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 144..6857
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..143
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 6855..7175
US-08-193-078B-8

Query Match 84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgagcttcctcagcgtt 20
|||||
DB 652 tccatgacgttcacgccggt 633

RESULT 12
US-08-223-305C-8/C
Sequence 8, Application US/08223305C
Patent No. 5851824
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
NUMBER OF SEQUENCES: 57
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ Version 1.5
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/223,305C
FILING DATE: April 4, 1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/868,354
FILING DATE: April 10, 1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206

FILED DATE: 15-AUG-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/620,250
FILING DATE: 30-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/482,384
FILING DATE: 20-FEB-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/603,751
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US89/01408
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/176,899
FILING DATE: 04-APR-1988
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 52516 (P519739)
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619)238-0999
TELEFAX: (619)238-0062
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 7175 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 144..6857
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..143
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 6855..7175
US-08-223-305C-8

Query Match 84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgcagctt 20
|||||
Db 652 tccatgacgttcctgcagctt 633

RESULT 13
US-08-149-097D-B/c
Sequence 8, Application US/08149097D
Patent No. 5874236
GENERAL INFORMATION:
APPLICANT: Harpold, Michael
APPLICANT: Ellis, Steven
APPLICANT: Williams, Mark
APPLICANT: Feldman, Daniel
APPLICANT: McCue, Ann
APPLICANT: Brenner, Robert
TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
TITLE OF INVENTION: METHODS
NUMBER OF SEQUENCES: 40
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brown, Martin, Haller & McClain
STREET: 1660 Union Street
CITY: San Diego
STATE: California
COUNTRY: USA
ZIP: 92101-2926
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/149,097D
FILING DATE: 05-NOV-1993
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/105,536
FILING DATE: 11-AUG-1993
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US92/06903
FILING DATE: 14-AUG-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/914,231
FILING DATE: 13-JUL-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/868,354
FILING DATE: 10-APR-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/745,206
FILING DATE: 15-AUG-1991
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/620,250
FILING DATE: 30-NOV-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/482,384
FILING DATE: 20-FEB-1990
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/603,751
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: WO PCT/US89/01408
FILING DATE: 04-APR-1989
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/176,899
FILING DATE: 04-APR-1988
ATTORNEY/AGENT INFORMATION:
NAME: Seidman, Stephanie L.
REGISTRATION NUMBER: 33,779
REFERENCE/DOCKET NUMBER: 6362-55038
TELECOMMUNICATION INFORMATION:
TELEPHONE: (619) 238-0999
TELEFAX: (619) 238-0062
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 7175 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: CDS
LOCATION: 144..6857
FEATURE:
NAME/KEY: 5'UTR
LOCATION: 1..143
FEATURE:
NAME/KEY: 3'UTR
LOCATION: 6855..7175
US-08-149-097D-8

Query Match 84.0%; Score 16.8; DB 3; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgcagctt 20
|||||
Db 652 tccatgacgttcctgcagctt 633

RESULT 14
US-08-949-386-8/C
; Sequence 8, Application US/08949386
; Patent No. 6090623
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Gillespie, Alison
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/949,386
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/290,012
; FILING DATE: 11-AUG-1994
; APPLICATION NUMBER: 08/149,097
; FILING DATE: 5-NOV-1993
; PRIOR APPLICATION DATA: 08/105,536
; FILING DATE: 11-AUG-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 519808
; TELEPHONE: (619) 238-0062
; TELEFAX: (619) 238-0999
; INFORMATION FOR SEQ. ID NO.: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7175 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 144..6857
; FEATURE:
; NAME/KEY: 5'UTR
; LOCATION: 1..143
; FEATURE:
; NAME/KEY: 3'UTR
; LOCATION: 6855..7175
; US-08-949-386-8

Query Match 84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgtctcctgacgtc 20
|||||
Db 652 tccatgacgttccacccgtt 633

RESULT 15

US-08-450-562-8/C
; Sequence 8, Application US/08450562
; Patent No. 6096514
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Gillespie, Alison
; APPLICANT: Feldman, Daniel
; APPLICANT: Brenner, Robert
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/450,562
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/404,950
; FILING DATE: 13-MAR-1995
; APPLICATION NUMBER: 08/336,257
; FILING DATE: 7-NOV-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/314,083
; FILING DATE: 28-SEPT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/311,363
; FILING DATE: 23-SEPT-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/290,012
; FILING DATE: 11-AUG-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/223,305
; FILING DATE: 4-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/193,078
; FILING DATE: 07-FEB-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/149,097
; FILING DATE: 5-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/105,536
; FILING DATE: 11-AUG-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/914,231
; FILING DATE: 13-JULY-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/868,354
; FILING DATE: 10-APR-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/06903
; FILING DATE: 14-AUG-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/745,206
; FILING DATE: 15-AUG-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/620,250
; FILING DATE: 30-NOV-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/603,751

;; FILING DATE: 08-NOV-1990
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 07/482,384
;; FILING DATE: 02-FEB-1990
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: PCT/US89/01408
;; FILING DATE: 04-APR-1989
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: 07/176,899
;; FILING DATE: 04-APR-1988
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Seidman, Stephanie L.
;; REGISTRATION NUMBER: 33,779
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: (619) 238-0999
;; TELEFAX: (619) 238-0062
;; INFORMATION FOR SEQ ID NO: 8:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 7175 base pairs
;; TYPE: nucleic acid
;; STRANDEDNESS: double
;; TOPOLOGY: linear
;; MOLECULE TYPE: DNA (genomic)
;; FEATURE:
;; NAME/KEY: CDS
;; LOCATION: 144..6857
;; FEATURE:
;; NAME/KEY: 5'UTR
;; LOCATION: 1..143
;; FEATURE:
;; NAME/KEY: 3'UTR
;; LOCATION: 6855..7175
;; US-08-450-562-8

Query Match 84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 7.6;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 tccatgacgttcctgacgtt 20
|||||
Db 652 tccatgacgttcacgacctt 633

Search completed: December 4, 2000, 21:08:06
Job time: 16804 sec

GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 21:09:55 ; Search time 99.31 Seconds
(without alignments)
75.655 Million cell updates/sec

Title: US-09-369-941-2
Perfect score: 20
Sequence: 1 tccatgacgtctctgacgtt 20

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 480022 segs, 187831343 residues
Total number of hits satisfying chosen parameters: 960044

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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3: /cgn2_2/gcgdata/geneseq/geneseq/NA1981.DAT.*
4: /cgn2_2/gcgdata/geneseq/geneseq/NA1983.DAT.*
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20: /cgn2_2/gcgdata/geneseq/geneseq/NA1999.DAT.*
21: /cgn2_2/gcgdata/geneseq/geneseq/NA2000.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	19 V60950	Unmethylated cytos
2	20	100.0	20	19 V47683	Unmethylated Cpg d
3	20	100.0	20	19 V27667	Immunostimulatory
4	20	100.0	20	20 Z31943	Cpg adjuvant oligo
5	20	100.0	20	20 Z41946	IL-12 secretion in
6	20	100.0	20	20 Z41949	IL-12 secretion in
7	20	100.0	20	20 Z28191	Chlamydia trachoma
8	20	100.0	20	20 X78802	HPV fusion protein
9	20	100.0	20	20 X88536	Cytosine-guanosine
10	20	100.0	20	20 V74237	Cpg-N motif S-ODN
11	20	100.0	20	20 V74242	Cpg-N motif O-ODN
12	20	100.0	20	20 V74244	Cpg-N motif SOS-OD

13	20	100.0	20	21 Z59004	Cpg motif for immu
14	20	100.0	20	21 Z59174	Inflammatory card
15	20	100.0	20	21 Z61010	Nucleotide sequenc
16	20	100.0	20	21 Z61012	Nucleotide sequenc
17	20	100.0	20	21 Z47601	Murine immune syst
18	20	100.0	20	21 Z47885	Immunostimulatory
19	20	100.0	20	21 Z47887	Immunostimulatory
20	20	100.0	20	21 Z48022	Immune remodeling
21	20	100.0	20	21 Z48025	Immune remodeling
22	20	100.0	44	20 V83723	Murine-specific Cpg
23	20	100.0	44	20 V83722	Murine-specific Cpg
24	20	100.0	44	20 V83726	Cpg-optimised gene
25	17	85.0	17	19 V52557	Unmethylated Cpg d
26	17	85.0	17	19 V27731	Immunostimulatory
27	17	85.0	17	20 Z41916	IL-12 secretion in
28	17	85.0	17	21 Z60984	Nucleotide sequenc
29	17	85.0	17	21 Z47653	Parasitic infectio
30	17	85.0	17	21 Z47859	Immunostimulatory
31	17	85.0	17	21 Z47992	Immune remodeling
32	16.8	84.0	20	18 T88792	Synthetic phosphor
33	16.8	84.0	20	19 V52567	Unmethylated Cpg d
34	16.8	84.0	20	19 V45995	Immune adjuvant Cpg
35	16.8	84.0	20	19 V45996	Immune adjuvant Cpg
36	16.8	84.0	20	19 V27708	Immunostimulatory
37	16.8	84.0	20	19 V27700	Immunostimulatory
38	16.8	84.0	20	19 V27651	Immunostimulatory
39	16.8	84.0	20	19 V27638	Immunostimulatory
40	16.8	84.0	20	20 Z41879	IL-12 secretion in
41	16.8	84.0	20	20 Z41919	IL-12 secretion in
42	16.8	84.0	20	20 Z41930	IL-12 secretion in
43	16.8	84.0	20	20 Z28190	Chlamydia trachoma
44	16.8	84.0	20	20 V72500	Cpg motif containi
45	16.8	84.0	20	20 V74261	Cpg-N motif oligon

ALIGNMENTS

RESULT 1	
ID V60950	Standard; DNA; 20 BP.
AC V60950;	
DT 14-DEC-1998	(first entry)
DE	Unmethylated cytosine-guanine dinucleotide containing oligonucleotide 1.
XX	
KW	ss; unmethylated Cpg dinucleotide; immune response; natural killer cell;
KW	Th2 response; Th1 response; Th1 cytokine; hepatitis B.
OS	Synthetic.
XX	
PN W09840100-A1.	
PD	17-SEP-1998.
XX	
PF 10-MAR-1998;	98WO-US04703.
XX	
PR 10-MAR-1997;	97US-0040376.
XX	
PA (OTTA-) OTTAWA CIVIC LOEB RES INST.	
PA (QIAG-) QIAGEN GMBH.	
XX (IOWA) UNIV IOWA RES FOUND.	
XX	
PI Davis HL, Krieg AM, Schorr J;	
XX	
DR WPI; 1998-520792/44.	
XX	
PT	Use of oligonucleotides containing an unethylated Cpg dinucleotide
PT	- useful as, e.g. adjuvant with antigen, or nucleic acid encoding
PT	antigen for inducing immune response in subject
XX	

PS Claim 14; Page 35; 67pp; English.
XX
CC Oligonucleotides containing at least 1 unmethylated Cpg dinucleotide
CC affect the immune response in a subject by activating natural killer
CC cells or redirecting a subject's immune response from a Th2 to a Th1
CC response by inducing monocyte and other cells to produce Th1 cytokines.
CC These nucleic acids containing at least 1 unmethylated Cpg can be used as
CC an adjuvant, specifically to induce an immune response against an
CC antigenic protein, and are used particularly for virally mediated
CC disorders, e.g. hepatitis B virus infection.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgagcttcctgacgt 20
Db 1 tccatgagcttcctgacgt 20
|||||

RESULT 2
ID V47683 standard; DNA; 20 BP.
V47683:
AC V47683:
XX
XX 20-NOV-1998 (first entry)
DT
XX Unmethylated Cpg dinucleotide 1826.
DE
XX Unmethylated Cpg dinucleotide; immune response; bacterial meningitis;
XX natural killer cell activation; NK cell; Th2 response; neonatal sepsis;
KW pulmonary disorder; asthma; environmentally induced alway disease;
KW bacterial infection; endotoxaemia; therapy; cystic fibrosis;
KW inflammatory bowel disease; ss.
XX
XX Synthetic.
OS
XX
XX WO9837919-A1.
PN
XX 03-SEP-1998.
PD
XX 25-FEB-1998; 98WO-US03678.
PF
XX 28-FEB-1997; 97US-0039405.
PR
XX (IOWA) UNIV IOWA RES FOUND.
PA
XX Kriegl AM, Schwartz DA;
PI
XX WPI; 1998-480941/41.
DR
XX
XX
PT Use of nucleic acids containing an unmethylated Cpg - for treating a
PT subject having or at risk of having an acute decrement in air flow
PT or inhibiting an inflammatory response
XX
XX
PS Claim 35; Page 27; 65pp; English.
XX
CC This sequence represents an unmethylated Cpg dinucleotide, and can be
CC used in the method of the invention. The method is for treating a subject
CC having, or at risk of having an acute decrement in air flow, comprising
CC administering a nucleic acid sequence containing at least one
CC unmethylated Cpg. The nucleic acid sequence containing an unmethylated Cpg
CC dinucleotide affect an immune response in a subject by activating natural
CC killer cells (NK) or redirecting a subject's immune response from a Th2
CC to a Th1 response by inducing monocyte and other cells to produce Th1
CC cytokines. They can be used to treat pulmonary disorders having an
CC immunologic component, such as asthma or environmentally induced alway
CC disease. They can also be used to treat diseases associated with
CC Gram-positive bacterial infections or endotoxaemia including bacterial

CC meningitis, neonatal sepsis, cystic fibrosis, inflammatory bowel disease
CC and liver cirrhosis, Gram-negative pneumonia, Gram-negative abdominal
CC abscess, hemorrhagic shock, disseminated intravascular coagulation, or
CC an inflammatory response to lipopolysaccharide.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 tccatgagcttcctgacgt 20
Db 1 tccatgagcttcctgacgt 20
|||||

RESULT 3
ID V27667 standard; DNA; 20 BP.
V27667:
AC V27667:
XX
XX 01-OCT-1998 (first entry)
DT
XX Immunostimulatory oligodeoxyribonucleotide of the invention.
DE
XX Immunostimulatory; oligodeoxyribonucleotide; ODN;
KW unmethylated Cpg dinucleotide; activate; lymphocyte; immune response;
KW Th2; Th1; cytokine; treatment; prevention; asthma; autoimmune disease;
KW desensitisation therapy; artificial adjuvant; antibody generation; ss.
XX
XX Synthetic.
OS
XX
XX WO9818810-A1.
PN
XX 07-MAY-1998.
PD
XX 30-OCT-1997; 97WO-US19791.
PF
XX 30-OCT-1996; 96US-0738652.
PR
XX (IOWA) UNIV IOWA RES FOUND.
PA
XX Kline JN, Kriegl AM;
PI
XX WPI; 1998-272127/24.
DR
XX
XX
PT New immunostimulatory nucleic acid molecules - which contain at
PT least one unmethylated Cpg dinucleotide, used for treating e.g.
PT tumours, infections or autoimmune disease
XX
XX
PS Claim 35; Page 84; 109pp; English.
XX
CC V27641-751 represent immunostimulatory oligodeoxyribonucleotides (ODNs)
CC of the invention. The ODNs contain at least one unmethylated Cpg
CC dinucleotide, and have the formula:
CC 5' N1X1CGX2N2 3', where at least one nucleotide separates consecutive
CC Cpgs, X1 is adenine, guanine, or thymine, X2 is cytosine or thymine, N1 is
CC does not contain a CCGG tetramer or more than one CCG or CGG trimer OR
CC 5' N1X1X2CGX3X4N 3', where at least one nucleotide separates consecutive
CC Cpgs, X1 and X2 are selected from GPT, GPG, GpA, ApT and ApA, X3and X4
CC are selected from TPT or CPT. N is any nucleotide and N1-N2 is 0-26
CC bases with the provision that N1 and N2 does not contain a CCGG tetramer
CC or more than one CCG or CGG trimer.
CC The ODNs activate lymphocytes in a subject and redirect a subject's
CC immune response from a Th2 to a Th1 (e.g. by inducing monocyte cells and
CC other cells to produce Th1 cytokines, including IL-12, IFN-gamma and
CC GM-CSF). The ODNs can be used to treat or prevent an asthmatic disorder,
CC autoimmune diseases, in desensitisation therapy, as an artificial
CC adjuvant during antibody generation in a mammal such as a mouse or a
CC human.

```
XX SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
Query Match
Best Local Similarity 100.0%; Score 20; DB 19; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 tccatgacgttcctgacgtt 20
    |||
Db 1 tccatgacgttcctgacgtt 20

RESULT 4
Z31943
XX ID 231943 standard; DNA; 20 BP.
XX AC 231943;
XX DT 26-JAN-2000 (first entry)
XX DE Cpg adjuvant oligo 1001.
XX KM Cpg adjuvant; vaccine; polyoxyethylene ether; polyoxyethylene ester;
XX KW antigen; infection; allergy; cancer; therapy; ss.
XX OS Synthetic.
XX PA WO9952549-A1.
XX PN 21-OCT-1999.
XX PD 29-MAR-1999; 99WO-EP02278.
XX PF 09-APR-1998; 98GB-0007805.
XX PR 25-SEP-1998; 98GB-0020956.
XX PA (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.
XX PI Friede M, Hermand P;
XX DR WPI; 1999-620290/53.
XX PT Vaccine to protect against infections, allergy and cancer
XX PS Claim 16; Page 32; 52pp; English.
XX SQ This sequence represents a Cpg adjuvant that can be used in the vaccine
CC composition of the invention. The vaccine comprises a polyoxyethylene
CC ether or ester (I), not in the form of a vesicle, pharmaceutically
CC acceptable excipient and an antigen (Ag) or antigenic composition. The
CC vaccine can be used to treat or prevent infections (by bacteria, viruses
CC or other parasites), allergy and cancer. (I), which are safe, easy to
CC sterilize and simple to administer, are powerful vaccine adjuvants, able
CC to induce a systemic immune response when administered (non-invasively)
CC to the mucosa. The response is at least as good as that from conventional
CC systemic injection. (I) are effective at low concentration, have low
CC reactogenicity and are well tolerated.
XX SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
Query Match
Best Local Similarity 100.0%; Score 20; DB 20; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 tccatgacgttcctgacgtt 20
    |||
Db 1 tccatgacgttcctgacgtt 20

RESULT 5
Z41946
```

```
ID Z41946 standard; DNA; 20 BP.
XX AC Z41946;
XX DT 24-JAN-2000 (first entry)
XX DE IL-12 secretion inducing Cpg oligonucleotide 91.
XX KM Cpg oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
XX KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
XX KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
XX KW antigen presenting cell; infection; allergic disease.
XX OS Synthetic.
XX PA WO9951259-A2.
XX PN 14-OCT-1999.
XX PD 02-APR-1999; 99WO-US07335.
XX PF 03-APR-1998; 98US-0080729.
XX PR (IOWA ) UNIV IOWA RES FOUND.
XX PA Krieg AM, Weiner G;
XX PI WPI; 1999-620169/53.
XX DR Novel synergistic combinations of immunostimulatory oligonucleotides
XX PT and immunopotentiating cytokines are useful for stimulating the immune
XX PT system
XX PS Example 8; Page 88; 91pp; English.
XX SQ Sequences Z41856-Z41949 are phosphorothioate Cpg oligonucleotides which
CC are used in the invention to induce interleukin-12 (IL-12) secretion
CC from human PBMC. The invention comprises stimulating an immune response
CC in a subject comprising administering to a subject exposed to an antigen,
CC an immunopotentiating cytokine and an immunostimulatory Cpg
CC oligonucleotide to induce a synergistic antigen specific immune response.
CC The methods are useful for treating cancer by stimulating an antigen
CC specific immune response against a cancer antigen. The methods can also
CC be used to treat neoplastic disorders in humans, including but not
CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
CC for treating infectious diseases, e.g. viral diseases such as HIV,
CC bacterial diseases, and fungal diseases. The methods may also be used to
CC treat allergic diseases, e.g. asthma. The methods and compositions may
CC also be applied to treat cancer and tumors in non human subjects,
CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
CC be treated and include leukaemia, haemangioepithelioma and bovine ocular
CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
CC caused by the bacterium Corynebacterium pseudotuberculosis, and
CC contagious lung tumour of sheep caused by jaagsiekte may also be treated.
CC Cpg oligonucleotides can be useful in activating B cells, NK cells, and
CC antigen presenting cells, such as monocytes and macrophages. Cpg
CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
CC be used as an adjuvant in conjunction with tumour antigens to protect
CC against a tumour challenge.
XX SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
Query Match
Best Local Similarity 100.0%; Score 20; DB 20; Length 20;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 tccatgacgttcctgacgtt 20
    |||
Db 1 tccatgacgttcctgacgtt 20
```

RESULT 6
 241949 ID 241949 standard; DNA: 20 BP.
 XX 241949;
 DT 24-JAN-2000 (first entry)
 XX IL-12 secretion inducing CpG oligonucleotide 94.
 DE
 XX CpG oligonucleotide; phosphorothioate; interleukin-12; IL-12; secretion;
 KW human PBMC; immune response; cancer; HIV; bacterial disease; asthma;
 KW neoplastic disorder; jaagsiekte; B cell; NK cell; ss; cytokine;
 KW antigen presenting cell; infection; allergic disease.
 XX
 OS Synthetic.
 XX
 PN MO9951259-A2.
 XX
 PD 14-OCT-1999.
 XX
 PF 02-APR-1999; 99WO-US07335.
 XX
 PR 03-APR-1998; 98US-0080729.
 XX
 PA (IOWA) UNIV IOWA RES FOUND.
 XX
 PI Kriegl AM, Weiner G;
 XX
 DR WPI: 1999-620169/53.
 XX
 PT Novel synergistic combinations of immunostimulatory oligonucleotides
 PT and immunopotentiating cytokines are useful for stimulating the immune
 PT system -
 XX
 PS Example 8; Page 89; 91pp; English.
 XX
 PS Sequences 241856-241949 are phosphorothioate CpG oligonucleotides which
 CC are used in the invention to induce interleukin-12 (IL-12) secretion
 CC from human PBMC. The invention comprises stimulating an immune response
 CC in a subject comprising administering to a subject exposed to an antigen,
 CC an immunopotentiating cytokine and an immunostimulatory CpG
 CC oligonucleotide to induce a synergistic antigen specific immune response.
 CC The methods are useful for treating cancer by stimulating an antigen
 CC specific immune response against a cancer antigen. The methods can also
 CC be used to treat neoplastic disorders in humans, including but not
 CC limited to: sarcoma, carcinoma, fibroma, lymphoma, melanoma,
 CC neuroblastoma, retinoblastoma, and glioma. The methods are also useful
 CC for treating infectious diseases, e.g. viral diseases such as HIV,
 CC bacterial diseases, and fungal diseases. The methods may also be used to
 CC treat allergic diseases, e.g. asthma. The methods and compositions may
 CC also be applied to treat cancer and tumours in non human subjects,
 CC e.g. cats and dogs. Neoplasias affecting agricultural livestock may also
 CC be treated and include leukaemia, haemangioendothelioma and bovine ocular
 CC neoplasia. Chronic, infectious, contagious diseases of sheep and goats
 CC caused by the bacterium *Corynebacterium pseudotuberculosis*, and
 CC contagious lung tumour of sheep caused by *jaagsiekte* may also be treated.
 CC CpG oligonucleotides can be useful in activating B cells, NK cells, and
 CC antigen presenting cells, such as monocytes and macrophages. CpG
 CC oligonucleotides enhance antibody dependent cellular cytotoxicity and can
 CC be used as an adjuvant in conjunction with tumour antigens to protect
 CC against a tumour challenge.
 XX
 XX Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match	100.0%	Score 20;	DB 20;	Length 20;
Best Local Similarity	100.0%	Pred. No. 0.25;		
Matches	20;	Conservative 0;	Mismatches 0;	Indels 0;
1	tccatgacgttcctcgaagtt	20		
1	tccatgacgttcctcgaagtt	20		

RESULT	7
ID	228191
XX	228191 standard: DNA; 20 BP.
XX	228191;
XX	20-DEC-1999 (first entry)
DT	
DE	Chlamydia trachomatis outer membrane protein gene-derived Cpg oligo 4
XX	
KM	Heart disease; inflammatory; autoimmune; cardiomyopathy; adjuvant;
KW	Cpg motif; vaccine; ds.
XX	
OS	Synthetic.
OS	Chlamydia trachomatis.
XX	
Key	Location/Qualifiers
FH	modified_base 1..20
FT	/tag= a
FT	/mod_base= OTHER
FT	/note="OTHER = phosphorothioate linkage"

PN	US5962636-A.
XX	
PD	05-OCT-1999.
XX	
PE	12-AUG-1998; 980S-0133774.
XX	
PR	12-AUG-1998; 980S-0133774.
XX	
PA	(AMGE-) AMGEN CANADA INC.
XX	
PI	Bechmaier K, Hessel AJ, Penninger JM, Neu N;
XX	
DR	WPI; 1999-589735/50.
XX	
PT	Peptides that induce or suppress inflammatory cardiomyopathy -
XX	
PS	Example 2; Column 25; 17pp; English.

This sequence represents DNA encoding Chlamydia trachomatis 60 kD outer
 membrane protein (OMP) gene-derived CpG oligonucleotide 4. This
 oligonucleotide contains a CpG motif. It was tested for its ability to
 act as an adjuvant for the M7A alpha peptide (Y42723), which can induce
 inflammatory cardiomyopathy (ICM) in mice. It was found to act as a
 potent immunostimulant, whereas a motif (Y28193) from the same
 source which did not contain a CpG motif (Y28193) was hardly effective as
 an adjuvant. Inflammatory cardiomyopathy peptides (Y42723, Y42725-Y42731)
 can be used with such an adjuvant and an excipient in a vaccine for
 decreasing ICM.

50 Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match	100.0%	Score 20;	DB 20;	length 20;
Best local Similarity	100.0%	Pred. No. 0	25;	
Matches	20;	Conservative	0;	Mismatches 0;
				Indels 0;
				Gaps 0;
QY	1	tccatgacgttcctgacgtt	20	
Db	1	tccatgacgttcctgacgtt	20	

RESULT	8
X78802	
ID	X78802 standard; DNA; 20 BP.
XX	
AC	X78802;
XX	
DT	06-SEP-1999 (first entry)
XX	

DE	HPV fusion protein Cpg oligonucleotide 1.
XX	
XX	Fusion protein; E6 protein; E7 protein; E6/E7; immunomodulator; tumour;
XX	immunological fusion partner; Cpg oligonucleotide; immune response;
KW	HPV antigen; prevention; treatment; primer; ss.
XX	
OS	Synthetic.
OS	Human papillomavirus.
XX	
PN	WO933868-A2.
PD	08-JUL-1999.
XX	
PF	18-DEC-1998; 98WO-EP08563.
XX	
PR	24-DEC-1997; 97GB-0027262.
XX	
PA	(SMIK) SMITHKLINE BEECHAM BIOLOGICALS.
PI	Dalemans WLJ, Gerard CMG;
XX	
DR	WPI; 1999-405485/34.
PT	
PT	Composition comprising an E6, E7 or E6/E7 fusion protein from HPV to
XX	induce immune response to HPV
XX	
PS	Claim 11; Page 36; 62pp; English.
XX	
CC	X78791-X78801 represent nucleic acid sequences which encode novel
CC	constructs comprising an E6 or E7 protein or E6/E7 fusion protein from
CC	HPV (represented in Y23375-Y23386). These constructs are optionally
CC	linked to an immunological fusion partner and an immunomodulatory Cpg
CC	oligonucleotide. The products of the invention can be used to induce an
CC	immune response in a patient to an HPV antigen. They can also be used
CC	for preventing or treating HPV induced tumours. This sequence represents
CC	a Cpg oligonucleotide which is used in the method of the invention.
XX	
SQ	Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
	Query Match 100.0%; Score 20; DB 20; Length 20;
	Best local Similarity 100.0%; Pred. NO. 0.25;
	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy	1 tccatgacgttcctgacgtt 20
Db	1 tccatgacgttcctgacgtt 20
RESULT 9	
X88536	
ID	X88536 standard; DNA; 20 BP.
XX	
AC	X88536;
XX	
DT	10-SEP-1999 (first entry)
XX	
DE	Cytosine-guanosine dinucleotide motif oligonucleotide #3.
XX	
KW	Cytosine-guanosine dinucleotide motif; Cpg; immunomodulation;
KW	unmethylated; vaccine; immunostimulation; immune response;
KW	T-independent type 1 antigen; T-independent type 2 antigen;
KW	polysaccharide conjugate antigen; ss.
XX	
OS	Synthetic.
XX	
PN	WO9933488-A2.
XX	
PD	08-JUL-1999.
XX	
PF	18-DEC-1998; 98WO-EP08562.
XX	
PR	24-DEC-1997; 97GB-0027262.

XX (SMK) SMITHKLINE BEECHAM BIOLOGICALS.
PA
XX
PI Dalemans WLU, Laferriere CAJ, Priels J;
XX
DR WPI: 1999-405369/34.
XX
PT A vaccine composition for inducing a immune response to
PT T-independent type 1 or type 2 antigen or polysaccharide conjugate
XX antigen
XX
PS Claim 6; Page 31; 35pp; English.
XX
CC The present invention describes a formulation (A) comprising a
CC cytosine-guanosine dinucleotide motif (Cpg) oligonucleotide and
CC T-independent type 1 or type 2 antigens or polysaccharide conjugate
CC antigen. The present sequence represent a specifically claimed Cpg
CC oligonucleotide. A vaccine composition comprising the formulation is
CC used for inducing a immune response to T-independent type 1 or type 2
CC antigen or polysaccharide conjugate antigen. The use of
CC immunostimulatory Cpg oligonucleotide acts as an adjuvant to
CC pneumococcal polysaccharides.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 tccatgagcttcctgaagtt 20
| | | | | | | | | | |
Db 1 tccatgagcttcctgaagtt 20

RESULT 10
V74237
ID V74237 standard; DNA; 20 BP.
XX
AC V74237;
XX
DT 15-MAR-1999 (first entry)
XX
DE Cpg-N motif S-ODN 1826 DNA.
XX
KW Cpg-N motif: immunostimulation; antigen; Cpg-S motif; immunisation; ODN;
KM viral antigen; bacterial antigen; parasite; therapeutic; growth factor;
KW toxin; tumour suppressor; cytokine; apoptotic protein; interferon;
KM hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.
XX
OS Synthetic.
XX
PN WO9852581-A1.

26-NOV-1998.
XX
PD 20-MAY-1998; 98MO-US10408.
XX
PE 20-MAY-1997; 97US-0047233.
XX
PR 20-MAY-1997; 97US-0047209.

(OTTA-) OTTAWA CIVIC HOSPITAL, LOEB RES INST.
PA (QIAG-) QIAGEN GMBH.
PA (IOWA-) UNIV IOWA RES FOUND.
XX
PI Davis HL, Krieg AM, Schorr J, Wu T;
XX
DR WPI: 1999-059712/05.
XX
PT Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -
PT for enhancing the immunostimulatory effect of an antigen or
PT enhancing the expression of a therapeutic polypeptide
XX

PS Example 1; Page 64; 109pp; English.

XX V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a

CC method for enhancing the immunostimulatory effect of an antigen encoded

CC by nucleic acid contained in a nucleic acid construct. The method

CC involves determining the Cpg-N and Cpg-S motifs present in the construct,

CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a

CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a

CC nucleic acid construct having enhanced immunostimulatory efficacy. The

CC method can be used for immunisation against viral antigens, e.g. from

CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a

CC parasite. They can also be used for expression of a therapeutic

CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,

CC apoptotic proteins, interferons, hormones, clotting factors, ligands and

CC receptors.

SO Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

QY Query Match 100.0%; Score 20; DB 20; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.25;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 tccatgacgttcctcgacgt 20

|||||

1 tccatgacgttcctcgacgt 20

RESULT 11

V74242

ID V74242 standard; DNA: 20 BP.

AC V74242;

XX 15-MAR-1999 (first entry)

DT Cpg-N motif O-ODN 2061 DNA.

XX Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;

KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;

KM toxin; tumour suppressor; cytokine; apoptotic protein; interferon;

KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.

XX Synthetic.

OS WO9852581-A1.

XX 26-NOV-1998.

PD 20-MAY-1998; 98WO-US10408.

PF 20-MAY-1998; 98WO-US10408.

XX 20-MAY-1997; 97US-0047233.

PR 20-MAY-1997; 97US-0047209.

XX (OTTA-) OTTAWA CIVIC HOSPITAL LOEB RES INST.

PA (QIAG-) QIAGEN GMBH.

PA (IOWA-) UNIV IOWA RES FOUND.

XX Davis HL, Kriegl AM, Schorr J, Wu T;

PI WPI; 1999-059712/05.

DR

XX Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -

PT for enhancing the immunostimulatory effect of an antigen or

PT enhancing the expression of a therapeutic polypeptide

XX Example 1; Page 64; 109pp; English.

PS V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a

CC method for enhancing the immunostimulatory effect of an antigen encoded

CC by nucleic acid contained in a nucleic acid construct. The method

CC involves determining the Cpg-N and Cpg-S motifs present in the construct,

CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a

CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a

CC nucleic acid construct having enhanced immunostimulatory efficacy. The

CC method can be used for immunisation against viral antigens, e.g. from

CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a

CC parasite. They can also be used for expression of a therapeutic

CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,

CC apoptotic proteins, interferons, hormones, clotting factors, ligands and

CC receptors.

SO Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

QY Query Match 100.0%; Score 20; DB 20; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.25;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 tccatgacgttcctcgacgt 20

|||||

1 tccatgacgttcctcgacgt 20

RESULT 12

V74244

ID V74244 standard; DNA: 20 BP.

AC V74244;

XX 15-MAR-1999 (first entry)

DT Cpg-N motif SOS-ODN 1980 DNA.

XX Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;

KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;

KM toxin; tumour suppressor; cytokine; apoptotic protein; interferon;

KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.

XX Synthetic.

OS WO9852581-A1.

XX 26-NOV-1998.

PD 20-MAY-1998; 98WO-US10408.

PF 20-MAY-1997; 97US-0047233.

PR 20-MAY-1997; 97US-0047209.

XX (OTTA-) OTTAWA CIVIC HOSPITAL LOEB RES INST.

PA (QIAG-) QIAGEN GMBH.

PA (IOWA-) UNIV IOWA RES FOUND.

XX Davis HL, Kriegl AM, Schorr J, Wu T;

PI WPI; 1999-059712/05.

DR

XX Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -

PT for enhancing the immunostimulatory effect of an antigen or

PT enhancing the expression of a therapeutic polypeptide

XX Example 1; Page 64; 109pp; English.

PS V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a

CC method for enhancing the immunostimulatory effect of an antigen encoded

CC by nucleic acid contained in a nucleic acid construct. The method

CC involves determining the Cpg-N and Cpg-S motifs present in the construct,

CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a

CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a

CC nucleic acid construct having enhanced immunostimulatory efficacy. The

CC method can be used for immunisation against viral antigens, e.g. from

CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a

CC parasite. They can also be used for expression of a therapeutic

CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,

CC apoptotic proteins, interferons, hormones, clotting factors, ligands and

CC receptors.

CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a

CC nucleic acid construct having enhanced immunostimulatory efficacy. The

CC method can be used for immunisation against viral antigens, e.g. from

CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a

CC parasite. They can also be used for expression of a therapeutic

CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,

CC apoptotic proteins, interferons, hormones, clotting factors, ligands and

CC receptors.

SO Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

QY Query Match 100.0%; Score 20; DB 20; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.25;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 tccatgacgttcctcgacgt 20

|||||

1 tccatgacgttcctcgacgt 20

RESULT 12

V74244

ID V74244 standard; DNA: 20 BP.

AC V74244;

XX 15-MAR-1999 (first entry)

DT Cpg-N motif SOS-ODN 1980 DNA.

XX Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation; ODN;

KW viral antigen; bacterial antigen; parasite; therapeutic; growth factor;

KM toxin; tumour suppressor; cytokine; apoptotic protein; interferon;

KW hormone; clotting factor; ligand; receptor; oligodeoxynucleotide; ss.

XX Synthetic.

OS WO9852581-A1.

XX 26-NOV-1998.

PD 20-MAY-1998; 98WO-US10408.

PF 20-MAY-1997; 97US-0047233.

PR 20-MAY-1997; 97US-0047209.

XX (OTTA-) OTTAWA CIVIC HOSPITAL LOEB RES INST.

PA (QIAG-) QIAGEN GMBH.

PA (IOWA-) UNIV IOWA RES FOUND.

XX Davis HL, Kriegl AM, Schorr J, Wu T;

PI WPI; 1999-059712/05.

DR

XX Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors -

PT for enhancing the immunostimulatory effect of an antigen or

PT enhancing the expression of a therapeutic polypeptide

XX Example 1; Page 64; 109pp; English.

PS V74237-V74253 are oligodeoxynucleotide (ODN) primers used to describe a

CC method for enhancing the immunostimulatory effect of an antigen encoded

CC by nucleic acid contained in a nucleic acid construct. The method

CC involves determining the Cpg-N and Cpg-S motifs present in the construct,

CC removing neutralising Cpg (Cpg-N) motifs and optionally inserting a

CC stimulatory Cpg (Cpg-S) motifs in the construct, thereby producing a

CC nucleic acid construct having enhanced immunostimulatory efficacy. The

CC method can be used for immunisation against viral antigens, e.g. from

CC hepatitis B virus (HBV), bacterial antigens or an antigen derived from a

CC parasite. They can also be used for expression of a therapeutic

CC polypeptide, e.g. growth factors, toxins, tumour suppressors, cytokines,

CC apoptotic proteins, interferons, hormones, clotting factors, ligands and

CC receptors.

CC	receptors.
XX	
SQ	Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
	Query Match 100.0%; Score 20; DB 20; Length 20;
	Best Local Similarity 100.0%; Pred. No. 0.25;
	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 tccatgacgtcctcgactt 20
Db	1 tccatgacgtcctcgactt 20
RESULT 13	
ID	Z99004 standard; DNA; 20 BP.
XX	Z99004;
AC	
XX	21-JUN-2000 (first entry)
DE	Cpg motif for immunostimulatory oligonucleotide 1826.
XX	
KW	Immunoprotective; vaccine; antigen; saponin adjuvant; immune response; immunostimulatory oligonucleotide; unmethylated CpG dinucleotide; mammal; human; animal; ss.
KX	
OS	Synthetic.
XX	
PN	WO200009159-A1.
PD	24-FEB-2000.
PE	06-AUG-1999; 99WO-US17956.
XX	
PR	10-AUG-1998; 98US-0095913. 08-APR-1999; 99US-0128608.
PA	(AQUI-) ACUIIA BIOPHARMACEUTICALS INC.
PI	Kensil CA:
DR	WPI: 2000-224181/19.
PT	A vaccine composition comprising an antigen, saponin adjuvant and immunostimulatory CpG oligonucleotide, useful for stimulating immunity and increasing immune responses -
PS	Claim 10; Page 19; 38pp; English.
XX	
CC	The invention relates to a vaccine composition comprising an antigen, a saponin adjuvant and an immunostimulatory oligonucleotide. The immunostimulatory oligonucleotide preferably comprises at least one unmethylated CpG dinucleotide. This sequence represents an example of the immunostimulatory oligonucleotide. The vaccine composition increases the immune response to the antigen when administered to a mammal, especially a human or animal. It further stimulates immunity and especially enhances antibody production to the antigen, preferably in a positive synergistic manner. It further enhances cell-mediated immunity. The immune adjuvant in particular can be used to increase the immune response to an antigen in an individual or a test system.
SQ	Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;
	Query Match 100.0%; Score 20; DB 21; Length 20;
	Best Local Similarity 100.0%; Pred. No. 0.25;
	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0
QY	1 tccatgacgttcctcgactt 20
Db	1 tccatgacgttcctcgactt 20

RESULT 14
 Z99174
 ID Z99174 standard; DNA: 20 BP.
 XX Z99174;
 AC Z99174;
 DT 21-JUN-2000 (first entry)
 XX
 DE Inflammatory cardiomyopathy immunostimulatory oligonucleotide #3.
 XX
 KW Cardiant; murine alpha myosin heavy chain; inflammatory myocarditis;
 KW autoimmune inflammatory cardiomyopathy; Chlamydia; antibody; vaccine;
 KW hybridization probe; immunostimulatory; ss.
 XX
 OS Synthetic.
 XX
 PN US6034230-A.
 XX
 PD 07-MAR-2000.
 XX
 PF 03-MAY-1999; 99US-0303862.
 XX
 PR 12-AUG-1998; 98US-0133774.
 XX
 PA (AMGE-) AMGEN CANADA INC.
 XX
 PI Neu N, Penninger JM, Bachmayer K, Hessel AJ;
 XX
 DR WPI: 2000-255712/22.
 XX
 PT DNA molecules encoding novel myocardial peptides used for inhibiting
 PT and inducing inflammatory cardiomyopathy in vivo -
 XX
 PS Disclosure: Column 17; 17pp; English.
 XX
 CC The invention relates to the isolation of sequences coding for peptide
 CC sequences derived from bacteria and viruses which may cause inflammatory
 CC cardiomyopathy. The peptide sequences are searched based on the sequence
 CC of the M7A peptides derived from the murine alpha myosin heavy chain
 CC polypeptide. The consensus sequence of the murine M7A-alpha/beta peptides
 CC (Y83813) was used to search the PIR public database for similar bacterial
 CC and viral sequences able to cause inflammatory cardiomyopathy. The screen
 CC isolated the peptides Y83814-Y83819 and their corresponding coding
 CC sequences Z99164-Z99169. The peptides encoded by the DNAs are used, alone
 CC or in conjunction with other therapeutics, for inducing or inhibiting
 CC inflammatory cardiomyopathy in vivo, where the cardiomyopathy is
 CC autoimmune inflammatory cardiomyopathy, and inflammatory cardiomyopathy
 CC caused by Chlamydia or other bacterial or viral infections that cause
 CC inflammatory cardiomyopathy. The oligonucleotides Z99172-Z99176 were
 CC shown to increase the immunogenicity of the immunostimulatory peptides
 CC when injected simultaneously. The peptides may also be used for
 CC increasing inflammatory myocarditis in a mammal. Antibodies against the
 CC peptides and the peptides themselves are used for measuring the risk of
 CC inflammatory cardiomyopathy in a mammal. The peptides may also be used
 CC in vaccines. Nucleic acids encoding the peptides may be used as
 CC hybridization probes, e.g. in diagnostic assays to test for the
 CC presence of Chlamydia DNA.
 XX
 XX
 SQ Sequence 20 BP: 3 A; 6 C; 4 G; 7 T; 0 other;
 XX
 Query Match 100.0%; Score 20; DB 21; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.25; Indels 0; Gaps 0;
 Matches 20; Conservative 0; Mismatches 0;
 QY 1 tccatgacgttcctgacgtt 20
 ||||||||||||||||||
 Db 1 tccatgacgttcctgacgtt 20

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261010
ID 261010 standard; DNA; 20 BP.
XX
AC 261010;
XX
DE 30-MAY-2000 (first entry)
XX
DE Nucleotide sequence of an immunostimulatory CpG oligonucleotide.
XX
KW Immunostimulatory; stereoisomer; CpG oligonucleotide; Th2; Th1; asthma;
KW allergic reaction; allergen; cancer antigen; cancer; immunoinhibitory;
KW inflammatory disease; inflammatory bowel disease; autoimmune disease;
KW gingivitis; psoriasis; sepsis; ss.
XX
OS Synthetic.
XX
PN WO200006588-A1.
XX
PD 10-FEB-2000.
XX
PF 27-JUL-1999; 99WO-US17100.
XX
PR 27-JUL-1998; 98US-0094370.
XX
PA (IOWA ) UNIV IOWA RES FOUND.
PA (CPGT-) CPG IMMUNOPHARMACEUTICALS INC.
XX
PI Kriegl AM;
XX
PI
XX
DR WPL; 2000-195254/17.
XX
PT Immunostimulatory and immunoinhibitory stereoisomers of CpG
XX oligonucleotides useful for immunotherapy of cancer -
XX
PS Disclosure: Page 12; 88pp; English.
XX
CC 260933-261015 represent immunostimulatory stereoisomers of CpG
CC oligonucleotides. The sequences are derived from generic nucleic
CC acid sequence, from which immunoinhibitory sequences may also be
CC derived. The immunostimulatory nucleic acids can be co-administered
CC with an antigen to induce an antigen-specific immune response. The
CC immunostimulatory nucleic acids can also be used in methods for
CC redirecting a subject's immune response from a Th2 to a Th1, for
CC treating asthma, for desensitizing a subject against the occurrence
CC of an allergic reaction in response to contact with an allergen, for
CC activating an immune cell, especially a lymphocyte or a dendritic cell
CC expressing a cancer antigen or for treating cancer. The immunoinhibitory
CC nucleic acid can be used to prevent an immune response, especially where
CC the immune response in the subject is excessive due to having received
CC an immune stimulating compound. The immunoinhibitory nucleic acid can
CC be used to treat a subject having or at risk of an inflammatory disease,
CC especially inflammatory bowel disease, autoimmune disease, gingivitis,
CC psoriasis and sepsis.
XX
SQ Sequence 20 BP; 3 A; 6 C; 4 G; 7 T; 0 other;

```

```

Query Match          100.0%; Score 20; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 0.25;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 tccatgacgttcctcgcagctt 20
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Db 1 tccatgacgttcctcgcagctt 20

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Search completed: December 4, 2000, 21:09:55
 Job time: 16673 sec

DEFINITION Sequence 8 from patent US 5846757.

ACCESSION AR063883
VERSION AR063883.1 GI:5993191
KEYWORDS
SOURCE
ORGANISM Unknown.

REFERENCE
AUTHORS Unclassified.

1 (bases 1 to 7175)

Harpold, M.M., Ellis, S.B., Williams, M.E., Feldman, D.H., McCue, A.F.
and Brenner, R.

TITLE Human calcium channel .alpha..sub.1. .alpha..sub.2. and .beta.
subunits and assays using them

JOURNAL Patent: US 5846757-A 8 08-DEC-1998;

FEATURES Location/Qualifiers

source

1..7175

/organism="unknown"

BASE COUNT 1415 a 2197 c 2168 g 1395 t

ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 7175;

Best Local Similarity 90.0%; Pred. No. 2.4e+02;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttccgtgacgtt 20

|||||

Db 652 TCCATGACGTTCCAGCCGTT 633

RESULT 15

AR067883/c

LOCUS AR067883 7175 bp DNA

DEFINITION Sequence 8 from patent US 5851824.

ACCESSION AR067883

VERSION AR067883.1 GI:5999105

KEYWORDS

SOURCE Unknown.

ORGANISM Unknown.

REFERENCE 1 (bases 1 to 7175)

AUTHORS Harpold, M.M., Ellis, S.B., Williams, M.E., Feldman, D.H., McCue, A.F.
and Brenner, R.

TITLE Human calcium channel .alpha.-1C/.alpha.-1D, .alpha.-2, .beta.-1,
and .gamma.subunits and cells expressing the DNA

JOURNAL Patent: US 5851824-A 8 22-DEC-1998;

FEATURES Location/Qualifiers

source

1..7175

/organism="unknown"

BASE COUNT 1415 a 2197 c 2168 g 1395 t

ORIGIN

Query Match

Best Local Similarity 84.0%; Score 16.8; DB 5; Length 7175;

Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttccgtgacgtt 20

|||||

Db 652 TCCATGACGTTCCAGCCGTT 633

Search completed: December 4, 2000, 20:47:28

Job time: 18088 sec

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source 1..5467
BASE COUNT 1105 a 1598 c 1659 g 1105 t
ORIGIN
Query Match 84.0%; Score 16.8; DB 5; Length 5467;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttctcgtgacgtt 20
Db 652 TCCATGACGTTCCAGCGGTT 633

RESULT 12
D86600/c
LOCUS D86600 7113 bp mRNA INV 08-MAR-2000
DEFINITION Loligo bleekeri mRNA for voltage-dependent calcium channel, complete cds.
ACCESSION D86600
VERSION D86600.2 GI:7209875
KEYWORDS voltage-dependent calcium channel; neural calcium channel
SOURCE Loligo bleekeri cDNA to mRNA.
ORGANISM Loligo bleekeri
Eukaryota; Metazoa; Mollusca; Cephalopoda; Coleoidea; Teuthoidea; Myopsida; Lolliginidae; Loligo.
REFERENCE 1 (sites)
AUTHORS Kimura,T., Shouno,O., Hirota,K., Saito,T., Matsumoto,G. and Sato,C.
TITLE Molecular cloning and characterization of a putative neural calcium channel alpha1-subunit from squid optic lobe
MEDLINE Biochem. Biophys. Res. Commun. 230 (1), 147-154 (1997)
79148591
REFERENCE 2 (bases 1 to 7113)
AUTHORS Kimura,T., Shouno,O., Sato,C., Hirota,K., Hanyu,Y., Saito,T. and Matsumoto,G.
TITLE Primary structure of a putative calcium channel alpha1-subunit from squid optic lobe
JOURNAL Unpublished (1996)
REFERENCE 3 (bases 1 to 7113)
AUTHORS Kimura,T.
TITLE Direct Submission
JOURNAL Submitted (22-JUL-1996) to the DDBJ/EMBL/GenBank databases. Tadashi Kimura, Electrotechnical Laboratory, Supermolecular Science Division; 1-1-4 Umezono, Tsukuba, Ibaraki 305, Japan (E-mail:eveda@etl.go.jp, Tel:029854508066616)
COMMENT On Mar 8, 2000 this sequence version replaced gi:1817549.
Sequence updated (27-Nov-1996) by: Tadashi Kimura
Sequence updated (06-Mar-2000).
FEATURES
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1..7113
/organism="Loligo bleekeri"
/db_xref="taxon:6617"
329..6619
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/codon_start=1
/product="voltage-dependent calcium channel"
/protein_id="BAA13136.2"
/db_xref="GI:7209875"
/translation="MNTSADTGGRDGDGYTHDGSLOFVAKKAATVSLPGMGSSTNRS
LFIFSEENFIRKAYKIIIEWGPFYEWLLTIANCIVLALEELHPNEDKTPLAQVLEA
TEFYLFQFCFVAKLIVAGLFGALHGKSYLRNVNIMDFVVTGFIISFPASNSFDL
RTLRAVRLPLKLVGPSLQVVLKSIIRAMAPLQVCLLVLAIFAIVIGLEFYT
GAFKACFIKPNDESNDLEYGDEDTIRPCANEGSGYHCRANIKACRFNAGPNYGIT
SEDNMGFAMLVFOCVTWEGHTOVLYYTDGADYNNIYFVPLVLGSEFFMLNLVIG
VLSEFAKERERVENRAFLRROQIERENGLWEICKAEVILDEERKKDDGTI
SDEKLRIIEGALGNARLAAQVKMKKENKELTIGDNDGDLGSLNVGSGFRG
LKNRAHGRCAGFWAERKHLRFTIRKCVKTQGYWFVILVFLNTLCVASEHYGQAEW
HTEFLYVMEFAFLALFMSEMLIKMYGLGVRLYFQSFNIFDCWTLVLSIIEVTSIAIK
DGSSFGISTLRLLRMFKVTRYWSSLRNLVLSLMSRSIVSLFLFLFLFLPAL
LGMOLFGGMNMFEEGRPPGHEDTFPALLTVFQILTGEDMNEVSGIRAGGIAGGG
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NSEEPAFGPKPMLPYSSMFIQPTNPVRFFCHVNLRYEDLFIIVLICASSALAA
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ICALVAFAGDAGAGNLTKISURVLRVLRPLKTKAVFVCDVNSLKNVSNIL
LIVYLLQFIFAVIAVOLFKEFFCYTDMKSNRECCQGYFDEYEDSKPRVKNRW
LRQDFHDVNMFAMLTFTVTTGEGWPMVLKNSMDSTSDMKPKPGYRMENAIYVVF
FVDFPFFVNFVALLIITFQEOGENELVDQDKQKCIERSIEAKPSCRYVPKNK
NSIKYKIMOVVYSPKFCVVMVIALNTLVLMKYVGSPTKEYKILLQNLNLAESVLT
IRICILKLAGFGIGVFRDRNMEDFIIVGSIIDVVTNVPSPASSPTGSRFLFRAA
RLVKLLRQGTITRLLLWTFELQSPKALPYVCLLIAMLFYIATGMQVFGNIRLDSKTS
INRHNFRFFYAVLLFRCATGESWQIIMSLGSPCDPESKMLDNSCGLDIATYIV
FVTFIFLCFLMLNLEFVIMDNFDYLTIRDTISLGHILDEYSRWAEYDPLAGSRVH
YTDMYEMLRMEPPVPGFGRNCPYLRACRKLIRNMPLKEDGTVHFSTLFLAVRESLS
IRMSSAEEMDKKDEEMREVILKRVMPVQKKIVDLVPPNHELNNGKLTGVKVGGLLI
AENWRAYKASONNSLTKTEKEELFWEDDIKEYREDEIREYDESDYKDDYQEYQ
YOEDCKDIYEDYNQVETLETESNKHVSTPLHSPCIAINGQQFQOQLOVSDSERPP
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RKDSTVQKSETASLOPSEKQDFSMGLRPEHAHAPRPGSGSSRGLNAOTVPLSP
VSPRSLPQSPLGSLASPSMHRSSPRRGLDVGAFASVNIQDQAHSTAEHDIRH
KRAYFHPCKPDDSLSVPTSPOMRGRSGRHRPPLQOQGVMSPLPGPTSRKKEPTF
YRSTSLNRSRSPSNLTPTSTLHQHEYYGSAGLTDRSRSPPTMTTPPKATRLKLPV
PSKSTLNLQAQTRPKNMPVMPSPVTPQPSKPSGFINFRLNASPTPIRVGTICQ
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BASE COUNT 2146 a 1383 c 1514 g 2070 t
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Query Match 84.0%; Score 16.8; DB 33; Length 7113;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttctcgtgacgtt 20
Db 747 TCCATGATGTTCCAGACGTT 728

RESULT 13
AR022380/c
LOCUS AR022380 7175 bp DNA PAT 05-DEC-1998
DEFINITION Sequence 8 from patent US 5792846.
ACCESSION AR022380
VERSION AR022380.1 GI:3976442
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 7175)
AUTHORS Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F. and Brenner,R.
TITLE Human calcium channel compositions and methods
JOURNAL Patent: US 5792846-A 8 11-AUG-1998;
FEATURES
Location/Qualifiers
source
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/organism="unknown"
BASE COUNT 1415 a 2197 c 2168 g 1395 t
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Query Match 84.0%; Score 16.8; DB 5; Length 7175;
Best Local Similarity 90.0%; Pred. No. 2.4e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttctcgtgacgtt 20
Db 652 TCCATGACGTTCCAGCGGTT 633

RESULT 14
AR063883/c
LOCUS AR063883 7175 bp DNA PAT 29-SEP-1999
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RESULT 7
LOCUS A90870 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent EP0855184.
ACCESSION A90870
VERSION A90870.1 GI:6739264
KEYWORDS
SOURCE
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg,K.P. and Lipford,G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
JOURNAL antigen especially for vaccination
PATENT: EP 0855184-A 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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source Location/Qualifiers
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ORIGIN

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Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 TCCATGACGTTCTGTGCT 20

RESULT 8
LOCUS A93512 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent WO9740163.
ACCESSION A93512
VERSION A93512.1 GI:6741731
KEYWORDS
SOURCE
ORGANISM unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Colpan,M. and Schorr,J.
TITLE NUCLEIC ACID VACCINATION FOR PARVOVIRAL INFECTIONS
JOURNAL Patent: WO 9740163-A 30-OCT-1997;
COLPAN METIN (DE); SCHORR JOACHIM (DE)
FEATURES
source Location/Qualifiers
1..20
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/db_xref="taxon:32644"
BASE COUNT 3 a 6 c 4 g 7 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 20;
Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 TCCATGACGTTCTGTGCT 20

RESULT 9
LOCUS A93521 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 14 from Patent WO9740163.
ACCESSION A93521
VERSION A93521.1 GI:6741738
KEYWORDS
SOURCE
ORGANISM unidentified.

ORGANISM unidentified
unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Colpan,M. and Schorr,J.
TITLE NUCLEIC ACID VACCINATION FOR PARVOVIRAL INFECTIONS
JOURNAL Patent: WO 9740163-A 30-OCT-1997;
COLPAN METIN (DE); SCHORR JOACHIM (DE)
FEATURES
source Location/Qualifiers
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/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT 3 a 6 c 4 g 7 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 20;
Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 1 TCCATGACGTTCTGTGCT 20

RESULT 10
LOCUS I12881/c 2470 bp DNA PAT 26-JUL-1995
DEFINITION Sequence 14 from patent US 5429921.
ACCESSION I12881
VERSION I12881.1 GI:910858
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 2470)
AUTHORS Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F.
and Brenner,R.
TITLE Assays for agonists and antagonists of recombinant human calcium
JOURNAL channels
PATENT: US 5429921-A 14 04-JUL-1995;
LOCATION/Qualifiers
source
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/organism="unknown"
BASE COUNT 483 a 722 c 754 g 511 t
ORIGIN

Query Match 84.0%; Score 16.8; DB 5; Length 2470;
Best Local Similarity 90.0%; Pred. No. 2.3e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 tccatgacgttcctgacgtt 20
|||||
Db 509 TCCATGACGTTCCAGCCGTT 490

RESULT 11
LOCUS I12880 5467 bp DNA PAT 26-JUL-1995
DEFINITION Sequence 12 from patent US 5429921.
ACCESSION I12880
VERSION I12880.1 GI:910857
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 5467)
AUTHORS Harpold,M.M., Ellis,S.B., Williams,M.E., Feldman,D.H., McCue,A.F.
and Brenner,R.
TITLE Assays for agonists and antagonists of recombinant human calcium
JOURNAL channels
PATENT: US 5429921-A 12 04-JUL-1995;
LOCATION/Qualifiers
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RESULT 3
AB01662S3/c
LOCUS AB01662S3 595 bp DNA ROD 14-APR-2000
DEFINITION Mus musculus gene for aldose reductase, exon 9.
ACCESSION AB016664
VERSION AB016664.1 GI:4586467
KEYWORDS aldose reductase.
SEGMENT 3 of 4
SOURCE Mus musculus (strain: BALB/c) DNA.
ORGANISM Mus musculus
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
1 (sites)
Li, H., Nobukuni, Y., Gui, T. and Yabe-Nishimura, C.
Characterization of genomic regions directing the cell-specific
expression of the mouse aldose reductase gene
Biochem. Biophys. Res. Commun. 255 (3), 759-764 (1999)
99160426
2 (bases 1 to 595)
Yabe-Nishimura, C. and Li, H.
Direct Submission
Submitted (31-JUL-1998) to the DDBJ/EMBL/GenBank databases. Chihiro
Yabe-Nishimura, Kyoto Prefectural University of Medicine,
Department of Pharmacology; Kawarachi-Hirokaji, Kamikyoku, Kyoto,
Kyoto 602-8566, Japan (E-mail: nchihiro@basic.kpu-m.ac.jp,
Tel: +81-75-251-5333, Fax: +81-75-251-5348)
FEATURES
source
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/strain="BALB/c"
/db_xref="taxon:10090"
276..358
/number=9
/product="aldose reductase"
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Query Match 87.0%; Score 17.4; DB 12; Length 595;
Best Local Similarity 94.7%; Pred. No. 1.1e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 ccatagcgttcctgcagctt 20
|||||
Db 215 CCATGACGTTCCAGAGCTT 197

RESULT 4
A89782
LOCUS A89782 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 4 from Patent WO9832462.
ACCESSION A89782
VERSION A89782.1 GI:6738296
KEYWORDS unidentified.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford, G.B. and Heeg, K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
JOURNAL OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
PATENT: WO 9832462-A 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
FEATURES
source
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/db_xref="taxon:32644"
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Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttcctgcagctt 20
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RESULT 5
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LOCUS A89783 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 5 from Patent WO9832462.
ACCESSION A89783
VERSION A89783.1 GI:6738297
KEYWORDS unidentified.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Lipford, G.B. and Heeg, K.
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND
JOURNAL OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION
PATENT: WO 9832462-A 30-JUL-1998;
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
FEATURES
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Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 1 TCCATGACGTTCTCTGATGCT 20

RESULT 6
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LOCUS A90869 20 bp DNA PAT 22-JAN-2000
DEFINITION Sequence 4 from Patent EP0855184.
ACCESSION A90869
VERSION A90869.1 GI:6739263
KEYWORDS unidentified.
SOURCE unidentified.
ORGANISM unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg, K.P. and Lipford, G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
JOURNAL antigen especially for vaccination
PATENT: EP 0855184-A 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
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Best Local Similarity 90.0%; Pred. No. 2.2e+02;
Matches 18; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 tccatgacgttcctgcagctt 20
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c 21 16.8 84.0 7713 32 RABBCBIII
c 22 16.8 84.0 30713 83 CELW05G11
c 23 16.8 84.0 69051 83 AC073171
c 24 16.8 84.0 124595 90 HS322E17
c 25 16.8 84.0 148871 78 AC025324
c 26 16.8 84.0 170595 78 AC025937
c 27 16.8 84.0 186526 69 AC008669
c 28 16.4 82.0 3694 12 AF224508
c 29 16 80.0 2271 7 AF135189
c 30 16 80.0 82686 7 AC004669
c 31 16 80.0 133137 10 AC007245
c 32 15.8 79.0 371 82 AC062158
c 33 15.8 79.0 615 81 AC048760
c 34 15.8 79.0 720 8 CNS01CTC
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c 36 15.8 79.0 762 81 AC051209
c 37 15.8 79.0 768 81 AC055639
c 38 15.8 79.0 777 84 AC075866
c 39 15.8 79.0 790 80 AC037305
c 40 15.8 79.0 799 81 AC058159
c 41 15.8 79.0 823 80 AC039484
c 42 15.8 79.0 873 81 AC049019
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ALIGNMENTS

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RESULT 1
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DEFINITION AF053408
ACCESSION AF053408
VERSION AF053408.1 GI:3746485
KEYWORDS Expression vector pMCG16-S.
ORGANISM Expression vector pMCG16-S.
SOURCE Expression vector pMCG16-S.
artificial sequence; vectors.
REFERENCE 1 (bases 1 to 4227)
AUTHORS Krieg,A.M., Wu,T., Weeratna,R., Efler,S.M., Love-Homan,L., Yang,L.,
Yi,A.K., Short,D. and Davis,H.L.
TITLE Sequence motifs in adenoviral DNA block immune activation by
stimulatory CpG motifs
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 95 (21), 12631-12636 (1998)
MEDLINE 9845422
2 (bases 1 to 4227)
AUTHORS Wu,T., Efler,S.M., Davis,H.L., Krieg,A.M. and Schorr,J.
DIRECT SUBMISSION
TITLE Submitted (12-MAR-1998) HGD, Loeb Health Research Institute, 725
Parkdale Ave., Ottawa, ON K1Y 4E9, Canada
JOURNAL Location/Qualifiers
FEATURES
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Db 2444 TCCATGACGTTCTCGACGTT 2463

RESULT 2
AF053409 4625 bp DNA circular SYN 17-OCT-1998
LOCUS Expression vector pMCG50-S, complete sequence.
DEFINITION AF053409
ACCESSION AF053409
VERSION AF053409.1 GI:3746487
KEYWORDS Expression vector pMCG50-S.
ORGANISM Expression vector pMCG50-S.
artificial sequence; vectors.
REFERENCE 1 (bases 1 to 4625)
AUTHORS Krieg,A.M., Wu,T., Weeratna,R., Efler,S.M., Love-Homan,L., Yang,L.,
Yi,A.K., Short,D. and Davis,H.L.
TITLE Stimulatory CpG motifs
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 95 (21), 12631-12636 (1998)
MEDLINE 98445422
2 (bases 1 to 4625)
AUTHORS Wu,T., Efler,S.M., Davis,H.L., Krieg,A.M. and Schorr,J.
DIRECT SUBMISSION
TITLE Submitted (12-MAR-1998) HGD, Loeb Health Research Institute, 725
Parkdale Ave., Ottawa, ON K1Y 4E9, Canada
JOURNAL Location/Qualifiers
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Best Local Similarity 100.0%; Pred. No. 4.4;
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QY 1 tccatgacgttcctgacgtt 20
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Db 2444 TCCATGACGTTCTCGACGTT 2463
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GenCore version 4.5
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OM nucleic - nucleic search, using sw model

Run on: December 4, 2000, 20:47:25 ; Search time 993.06 Seconds
(without alignments)
87.962 Million cell updates/sec

Title: US-09-369-941-2
Perfect score: 20
Sequence: 1 tccatgacgttcctgacgtt 20

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 1033670 seqs, 2183789903 residues
Total number of hits satisfying chosen parameters: 2067340

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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94: gb_vil2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	20	100.0	4625	13 AF053409	AF053409 Expressio
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4	16.8	84.0	20	5 A89782	A89782 Sequence 4
5	16.8	84.0	20	5 A89783	A89783 Sequence 5
6	16.8	84.0	20	5 A90869	A90869 Sequence 4
7	16.8	84.0	20	5 A90870	A90870 Sequence 5
8	16.8	84.0	20	5 A93512	A93512 Sequence 5
9	16.8	84.0	20	5 A93521	A93521 Sequence 14
C 10	16.8	84.0	2470	5 I12881	I12881 Sequence 14
C 11	16.8	84.0	5467	5 I12880	I12880 Sequence 12
C 12	16.8	84.0	7113	33 D86600	D86600 Loligo blea